(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification⁷: C07H 21/02, C12Q 1/68, C12P 21/00, 19/34, A01N 43/04, G06F 17/00

(21) International Application Number: PCT/US00/34758

(22) International Filing Date:
21 December 2000 (21.12.2000)

(25) Filing Language: English

(26) Publication Language: English

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(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:
— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: DRUG TARGET ISOGENES: POLYMORPHISMS IN THE NEUROPEPTIDE Y GENE

(57) Abstract: Polynucleotides comprising one or more of 8 novel single nucleotide polymorphisms in the human Neuropeptide Y (NPY) gene are described. Compositions and methods for detecting one or more of these polymorphisms are also disclosed. In addition, various genotypes and haplotypes for the NPY gene that exist in the population are described.

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DRUG TARGET ISOGENES:

POLYMORPHISMS IN THE NEUROPEPTIDE Y GENE

RELATED APPLICATIONS

5   This application claims the benefit of U.S. Provisional Application Serial No. 60/171,414 filed on December 21, 1999.

FIELD OF THE INVENTION

This invention relates to variation in genes that encode pharmaceutically important proteins.
10   In particular, this invention provides genetic variants of the human Neuropeptide Y (NPY) gene and methods for identifying which variant(s) of this gene is/are possessed by an individual.

BACKGROUND OF THE INVENTION

Current methods for identifying pharmaceuticals to treat disease often start by identifying,
15   cloning, and expressing an important target protein related to the disease. A determination of whether an agonist or antagonist is needed to produce an effect that may benefit a patient with the disease is then made. Then, vast numbers of compounds are screened against the target protein to find new potential drugs. The desired outcome of this process is a drug that is specific for the target, thereby reducing the incidence of the undesired side effects usually caused by a compound's activity
20   at non-intended targets.

What this approach fails to consider, however, is that natural variability exists in any and every population with respect to a particular protein. A target protein currently used to screen drugs typically is expressed by a gene cloned from an individual who was arbitrarily selected. However, the nucleotide sequence of a particular gene may vary tremendously among individuals. Subtle
25   alteration(s) in the primary nucleotide sequence of a gene encoding a target protein may be manifested as significant variation in expression of or in the structure and/or function of the protein. Such alterations may explain the relatively high degree of uncertainty inherent in treatment of individuals with drugs whose design is based upon a single representative example of the target. For example, it is well-established that some classes of drugs frequently have lower efficacy in some
30   individuals than others, which means such individuals and their physicians must weigh the possible benefit of a larger dosage against a greater risk of side effects. In addition, variable information on the biological function or effects of a particular protein may be due to different scientists unknowingly studying different isoforms of the gene encoding the protein. Thus, information on the type and frequency of genomic variation that exists for pharmaceutically important proteins would be
35   useful.

The organization of single nucleotide variations (polymorphisms) in the primary sequence of a gene into one of the limited number of combinations that exist as units of inheritance is termed a

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haplotype. Each haplotype therefore contains significantly more information than individual unorganized polymorphisms. Haplotypes provide an accurate measurement of the genomic variation in the two chromosomes of an individual and can be used to identify and distinguish between isoforms of genes coding for drug targets.

5   Many diseases are associated with specific variations in gene sequences. However while there are examples in which individual polymorphisms act as genetic markers for a particular phenotype, in many other cases an individual polymorphism may be found in a variety of genomic backgrounds, i.e., haplotypes, and therefore shows no definitive coupling between the polymorphism and the causative site for the phenotype (Clark AG et al. 1998 *Am J Hum Genet* 63:595-612;

10   Ulbrecht M et al. 2000 *Am J Respir Crit Care Med* 161: 469-74). Because a haplotype represents the variation across each form of a gene, it is superior to a SNP, or even multiple unordered SNPs in the gene, as a tool to determine whether a particular gene is involved in a particular phenotype. For example, analysis of the association between a particular phenotype and each observed haplotype for a gene suspected to be associated with that particular phenotype permits ranking of each haplotype

15   by its statistical power of prediction for the phenotype. Haplotypes found to be strongly associated with the phenotype can then have that positive association confirmed by alternative methods to minimize false positives. Conversely, if no observed haplotypes for that gene show association with the phenotype of interest, then it may be inferred that variation in the gene has little, if any, involvement with that phenotype. Thus, the ability to determine readily whether a gene is involved

20   in the cause or symptoms of a particular disease, or in the response to a particular drug, will be one useful application for haplotypes in development of diagnostics and drugs.

One possible drug target for the treatment of atherosclerosis, obesity, psychological disorders and alcoholism is the Neuropeptide Y (NPY) gene or its encoded product. NPY is a 36-amino acid vasoactive peptide widely distributed throughout the central and peripheral nervous system (Ito, H. et

25   al., Clin. Nephrol. 51:272-279, 1999; Karvonen, M.K. et al., Nat. Med. 4:1434-1437, 1998; Ann. Med 30:508-510, 1998). NPY has a vasocontrictor effect and enhances the effect of various agonists, including noradrenaline and angiotensin II (Pedrazzini et al., Curr Opin Nephrol Hypertens 1993 2(1):106-13). NPY plays an important role in the contraction of heart muscle by inducing an increase in the levels of cytosolic and nuclear calcium in heart and vascular smooth muscle cells

30   (Jacques et al., Can J. Physiol Pjarmacol 2000 78(2):162-72). NPY is a potent orexigenic agent and is believed to be involved in the regulation of eating behavior (Kokot and Ficek., Miner Electrolyte Metab 1999 25(4-6):303-5). It is also involved in the regulation of neuronal activity both under physiological conditions and during pathological hyperactivity such as seizures. Experimental models and tissue from epilepsy sufferers suggest that NPY-mediated neurotrasmission is altered in

35   seizures (Vezzani et al. 1999 22(1):25-30).

The Neuropeptide Y gene is located on chromosome 7pter-q22 and contains 3 exons that encode a 97 amino acid protein. Reference sequences for the NPY gene (reverse complement of a

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portion of GenBank Accession No: AC004485.1; SEQ ID NO:1), coding sequence (GenBank Accession No. NM_000905.1), and protein are shown in Figures 1, 2 and 3, respectively.

Several single nucleotide polymorphisms (SNPs) in the NPY gene have been reported in the NCBI SNP Database which correspond to the following polymorphisms in Figure 1: a cytosine or
5 adenine at a position corresponding to nucleotide 155413 (Ref SNP ID# 16146), a guanine or thymine at a position corresponding to nucleotide 155420 (Ref SNP ID# 16145), a guanine or adenine at a position corresponding to nucleotide 155562 (Ref SNP ID# 16144), a cytosine or thymine at a position corresponding to nucleotide 156019 (Ref SNP ID#16143), a guanine or adenine at a position corresponding to nucleotide 156039 (Ref SNP ID# 16478), an adenine or guanine at a
10 position corresponding to nucleotide 156101 (Ref SNP ID# 16142), a thymine or cytosine at a position corresponding to nucleotide 156310 (Ref SNP ID#16139), a guanine or adenine at a position corresponding to nucleotide 156374 (Ref SNP ID# 5572), an adenine or guanine at a position corresponding to nucleotide 162782 (Ref SNP ID#5576), an adenine or guanine at a position corresponding to nucleotide 162917 (Ref SNP ID# 16475) and a thymine or cytosine at a position
15 corresponding to nucleotide 162948 (Ref SNP ID# 16126).

The polymorphism of thymine or cytosine at nucleotide position 156310 in Fig. 1 results in an amino acid variation of leucine (Leu) or proline (Pro) at amino acid position 7 in Figure 3. An association has been identified between the polymorphism producing the Pro7 variant and higher serum levels of total and LDL cholesterol in obese Finnish and Dutch subjects (Karvonen et al.,
20 supra). Nishanen et al. (J. Clin. Endocrinol Metab 2000 86(6):2266-9) also reported that subjects having Pro7 had higher serum cholesterol and apolipoprotein B levels, and are at a higher risk for carotid artherosclerosis than individuals with the Leu7 variant. Recent studies on middle-aged men from Eastern Finland have shown that the Leu7Pro variation is associated with alcohol consumption. Individuals with Pro7 showed a higher average alcohol consumption than individuals with Leu7
25 (Kauhanen et al., Am. J. Med. Genet 2000 93(2):117-21). Because of the potential for polymorphisms in the NPY gene to affect the expression and function of the encoded protein, it would be useful to determine whether additional polymorphisms exist in the NPY gene, as well as how such polymorphisms are combined in different copies of the gene. Such information would be useful for studying the biological function of NPY as well as in identifying drugs targeting this
30 protein for the treatment of disorders related to its abnormal expression or function.

SUMMARY OF THE INVENTION

Accordingly, the inventors herein have discovered 8 novel polymorphic sites in the NPY gene. These polymorphic sites (PS) correspond to the following nucleotide positions in the reverse
35 complement of the indicated GenBank Accession Number: 154224 (PS1), 154841 (PS2), 155063 (PS3), 155119 (PS4), 155412 (PS5), 155954 (PS9), 156117 (PS13) and 156259 (PS14) in AC004485.1. The polymorphisms at these sites are thymine or guanine at PS1, thymine or cytosine

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at PS2, guanine or thymine at PS3, cytosine or adenine at PS4, guanine or thymine at PS5, thymine
or cytosine at PS9, adenine or guanine at PS13 and adenine or cytosine at PS14. In addition, the
inventors have determined the identity of the alternative nucleotides present at these sites, as well as
at the previously identified sites at nucleotides 155413 (PS6), 155420 (PS7), 155562 (PS8),156019

5   (PS10), 156039 (PS11), 156101 (PS12), 156310 (PS15), 156374 (PS16), 162782 (PS17), 162917
(PS18) and 162948 (PS19), in a human reference population of 79 unrelated individuals self-
identified as belonging to one of four major population groups: African descent, Asian, Caucasian
and Hispanic/Latino. It is believed that NPY-encoding polynucleotides containing one or more of the
novel polymorphic sites reported herein will be useful in studying the expression and biological

10   function of NPY, as well as in developing drugs targeting this protein. In addition, information on
the combination of polymorphisms in the NPY gene may have diagnostic and forensic applications.

     Thus, in one embodiment, the invention provides an isolated polynucleotide comprising a
nucleotide sequence which is a polymorphic variant of a reference sequence for the NPY gene or a
fragment thereof. The reference sequence comprises SEQ ID NO:1 and the polymorphic variant

15   comprises at least one polymorphism selected from the group consisting of guanine at PS1, cytosine
at PS2, thymine at PS3, adenine at PS4, thymine at PS5, cytosine at PS9, guanine at PS13 and
cytosine at PS14. In a preferred embodiment, the polymorphic variant comprises one or more
additional polymorphisms selected from the group consisting of adenine at PS6, thymine at PS7,
adenine at PS8, thymine at PS10, adenine at PS11; guanine at PS12, cytosine at PS15, adenine at

20   PS16, guanine at PS17, guanine at PS18 and cytosine at PS19. A particularly preferred polymorphic
variant is a naturally-occurring isoform (also referred to herein as an "isogene") of the NPY gene. A
NPY isogene of the invention comprises thymine or guanine at PS1, thymine or cytosine at PS2,
guanine or thymine at PS3, cytosine or adenine at PS4, guanine or thymine at PS5, cytosine or
adenine at PS6, guanine or thymine at PS7, guanine or adenine at PS8, thymine or cytosine at PS9,

25   cytosine or thymine at PS10, guanine or adenine at PS11, adenine or guanine at PS12, adenine or
guanine at PS13, adenine or cytosine at PS14, thymine or cytosine at PS15, guanine or adenine at
PS16, adenine or guanine at PS17, adenine or guanine at PS18 and thymine or cytosine at PS19. The
invention also provides a collection of NPY isogenes, referred to herein as a NPY genome anthology.

     A NPY isogene may be defined by the combination and order of these polymorphisms in the
30   isogene, which is referred to herein as a NPY haplotype. Thus, the invention also provides data on
the number of different NPY haplotypes found in the above four population groups. This haplotype
data is useful in methods for deriving a NPY haplotype from an individual's genotype for the NPY
gene and for determining an association between a NPY haplotype and a particular trait.

     Polynucleotides complementary to these NPY genomic DNA variants are also provided by
35   the invention.

     In other embodiments, the invention provides a recombinant expression vector comprising
one of the polymorphic genomic variants operably linked to expression regulatory elements as well

as a recombinant host cell transformed or transfected with the expression vector. The recombinant vector and host cell may be used to express NPY for protein structure analysis and drug binding studies.

In other embodiments, the invention provides methods, compositions, and kits for haplotyping and/or genotyping the NPY gene in an individual. The methods involve identifying the nucleotide or nucleotide pair present at one or more polymorphic sites selected from PS1-PS5, PS9, PS13 and PS14 in one or both copies of the NPY gene from the individual. The compositions contain oligonucleotide probes and primers designed to specifically hybridize to one or more target regions containing, or that are adjacent to, a polymorphic site. The methods and compositions for establishing the genotype or haplotype of an individual at the novel polymorphic sites described herein are useful for studying population diversity, anthropological lineage, the significance of diversity and lineage at the phenotypic level, paternity testing, forensic applications, and for identifying associations between the NPY genetic variation and a trait such as level of drug response or susceptibility to disease.

In yet another embodiment, the invention provides a method for identifying an association between a genotype or haplotype and a trait. In preferred embodiments, the trait is susceptibility to a disease, severity of a disease, the staging of a disease or response to a drug. Such methods have applicability in developing diagnostic tests and therapeutic treatments for atherosclerosis, obesity, psychological disorders and alcoholism.

The present invention also provides nonhuman transgenic animals comprising one of the NPY polymorphic genomic variants described herein and methods for producing such animals. The transgenic animals are useful for studying expression of the NPY isogenes *in vivo*, for *in vivo* screening and testing of drugs targeted against NPY protein, and for testing the efficacy of therapeutic agents and compounds for atherosclerosis, obesity, psychological disorders and alcoholism in a biological system.

The present invention also provides a computer system for storing and displaying polymorphism data determined for the NPY gene. The computer system comprises a computer processing unit; a display; and a database containing the polymorphism data. The polymorphism data includes the polymorphisms, the genotypes and the haplotypes identified for the NPY gene in a reference population. In a preferred embodiment, the computer system is capable of producing a display showing NPY haplotypes organized according to their evolutionary relationships.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 illustrates a reference sequence for the NPY gene that is the reverse complement of part of Genbank Accession Number AC004485.1 (contiguous lines; SEQ ID NO:1), with the start and stop positions of each region of coding sequence indicated below the sequence by the numbers within the brackets and the polymorphic sites and polymorphisms identified by Applicants in a

reference population indicated by the variant nucleotide positioned below the polymorphic site in the sequence. SEQ ID NO:89 is equivalent to Figure 1, with the two allelic variants of each polymorphic site indicated by the appropriate nucleotide symbol (r= g or a, y= t or c, m= a or c, k= g or t, s= g or c, and w= a or t; WIPO standard ST.25).

5   Figure 2 illustrates a reference sequence for the NPY coding sequence (contiguous lines; SEQ ID NO:2), with the polymorphic site and polymorphism identified by Applicants in a reference population indicated by the variant nucleotide positioned below the polymorphic site in the sequence.

        Figure 3 illustrates a reference sequence for the NPY protein (contiguous lines; SEQ ID NO:3), with the variant amino acid(s) caused by the polymorphism(s) of Fig. 2 positioned below the
10   polymorphic site in the sequence.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

        The present invention is based on the discovery of novel variants of the NPY gene. As described in more detail below, the inventors herein discovered 8 novel polymorphic sites by
15   characterizing the NPY gene found in genomic DNAs isolated from an Index Repository that contains immortalized cell lines from one chimpanzee and 93 human individuals. The human individuals included a reference population of 79 unrelated individuals self-identified as belonging to one of four major population groups: Caucasian (22 individuals), African descent (20 individuals) Asian (20 individuals) Hispanic/Latino (17 individuals). To the extent possible, the members of this
20   reference population were organized into population subgroups by the self-identified ethnogeographic origin of their four grandparents as shown in Table 1 below.

Table 1. Population Groups in the Index Repository

Population Group	Population Subgroup	No. of Individuals
African descent		20
	Sierra Leone	1
Asian		20
	Burma	1
	China	3
	Japan	6
	Korea	1
	Philippines	5
	Vietnam	4
Caucasian		22
	British Isles	3
	British Isles/Central	4
	British Isles/Eastern	1
	Central/Eastern	1
	Eastern	3
	Central/Mediterranean	1
	Mediterranean	2
	Scandinavian	2
Hispanic/Latino		17
	Caribbean	7
	Caribbean (Spanish Descent)	2
	Central American (Spanish Descent)	1
	Mexican American	4
	South American (Spanish Descent)	3

In addition, the Index Repository contains three unrelated indigenous American Indians (one from each of North, Central and South America), one three-generation Caucasian family (from the

5   CEPH Utah cohort) and one two-generation African-American family.

Using the NPY genotypes identified in the Index Repository and the methodology described in the Examples below, the inventors herein also determined the haplotypes found on each chromosome for most human members of this repository. The NPY genotypes and haplotypes found in the repository include those shown in Tables 4 and 5, respectively. The polymorphism and

10   haplotype data disclosed herein are useful for studying population diversity, anthropological lineage, the significance of diversity and lineage at the phenotypic level, paternity testing, forensic applications, and for identifying associations between the NPY genetic variation and a trait such as level of drug response or susceptibility to disease.

In the context of this disclosure, the following terms shall be defined as follows unless

15   otherwise indicated:

Allele - A particular form of a genetic locus, distinguished from other forms by its particular nucleotide sequence.

Candidate Gene – A gene which is hypothesized to be responsible for a disease, condition, or the response to a treatment, or to be correlated with one of these.

20   Gene - A segment of DNA that contains all the information for the regulated biosynthesis of

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an RNA product, including promoters, exons, introns, and other untranslated regions that control expression.

Genotype – An unphased 5′ to 3′ sequence of nucleotide pair(s) found at one or more polymorphic sites in a locus on a pair of homologous chromosomes in an individual. As used herein,

5   genotype includes a full-genotype and/or a sub-genotype as described below.

Full-genotype – The unphased 5′ to 3′ sequence of nucleotide pairs found at all known polymorphic sites in a locus on a pair of homologous chromosomes in a single individual.

Sub-genotype – The unphased 5′ to 3′ sequence of nucleotides seen at a subset of the known polymorphic sites in a locus on a pair of homologous chromosomes in a single individual.

10   Genotyping – A process for determining a genotype of an individual.

Haplotype – A 5′ to 3′ sequence of nucleotides found at one or more polymorphic sites in a locus on a single chromosome from a single individual. As used herein, haplotype includes a full-haplotype and/or a sub-haplotype as described below.

Full-haplotype – The 5′ to 3′ sequence of nucleotides found at all known polymorphic sites

15   in a locus on a single chromosome from a single individual.

Sub-haplotype – The 5′ to 3′ sequence of nucleotides seen at a subset of the known polymorphic sites in a locus on a single chromosome from a single individual.

Haplotype pair – The two haplotypes found for a locus in a single individual.

Haplotyping – A process for determining one or more haplotypes in an individual and

20   includes use of family pedigrees, molecular techniques and/or statistical inference.

Haplotype data - Information concerning one or more of the following for a specific gene: a listing of the haplotype pairs in each individual in a population; a listing of the different haplotypes in a population; frequency of each haplotype in that or other populations, and any known associations between one or more haplotypes and a trait.

25   Isoform – A particular form of a gene, mRNA, cDNA or the protein encoded thereby, distinguished from other forms by its particular sequence and/or structure.

Isogene – One of the isoforms of a gene found in a population. An isogene contains all of the polymorphisms present in the particular isoform of the gene.

Isolated – As applied to a biological molecule such as RNA, DNA, oligonucleotide, or

30   protein, isolated means the molecule is substantially free of other biological molecules such as nucleic acids, proteins, lipids, carbohydrates, or other material such as cellular debris and growth media. Generally, the term "isolated" is not intended to refer to a complete absence of such material or to absence of water, buffers, or salts, unless they are present in amounts that substantially interfere with the methods of the present invention.

35   Locus - A location on a chromosome or DNA molecule corresponding to a gene or a physical or phenotypic feature.

Naturally-occurring – A term used to designate that the object it is applied to, e.g.,

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naturally-occurring polynucleotide or polypeptide, can be isolated from a source in nature and which has not been intentionally modified by man.

Nucleotide pair – The nucleotides found at a polymorphic site on the two copies of a chromosome from an individual.

5 **Phased** – As applied to a sequence of nucleotide pairs for two or more polymorphic sites in a locus, phased means the combination of nucleotides present at those polymorphic sites on a single copy of the locus is known.

Polymorphic site (PS) – A position within a locus at which at least two alternative sequences are found in a population, the most frequent of which has a frequency of no more than 99%.

10 **Polymorphic variant** – A gene, mRNA, cDNA, polypeptide or peptide whose nucleotide or amino acid sequence varies from a reference sequence due to the presence of a polymorphism in the gene.

Polymorphism – The sequence variation observed in an individual at a polymorphic site. Polymorphisms include nucleotide substitutions, insertions, deletions and microsatellites and may,

15 but need not, result in detectable differences in gene expression or protein function.

Polymorphism data – Information concerning one or more of the following for a specific gene: location of polymorphic sites; sequence variation at those sites; frequency of polymorphisms in one or more populations; the different genotypes and/or haplotypes determined for the gene; frequency of one or more of these genotypes and/or haplotypes in one or more populations; any

20 known association(s) between a trait and a genotype or a haplotype for the gene.

Polymorphism Database – A collection of polymorphism data arranged in a systematic or methodical way and capable of being individually accessed by electronic or other means.

Polynucleotide – A nucleic acid molecule comprised of single-stranded RNA or DNA or comprised of complementary, double-stranded DNA.

25 **Population Group** – A group of individuals sharing a common ethnogeographic origin.

Reference Population – A group of subjects or individuals who are predicted to be representative of the genetic variation found in the general population. Typically, the reference population represents the genetic variation in the population at a certainty level of at least 85%, preferably at least 90%, more preferably at least 95% and even more preferably at least 99%.

30 **Single Nucleotide Polymorphism (SNP)** – Typically, the specific pair of nucleotides observed at a single polymorphic site. In rare cases, three or four nucleotides may be found.

Subject – A human individual whose genotypes or haplotypes or response to treatment or disease state are to be determined.

Treatment - A stimulus administered internally or externally to a subject.

35 **Unphased** – As applied to a sequence of nucleotide pairs for two or more polymorphic sites in a locus, unphased means the combination of nucleotides present at those polymorphic sites on a single copy of the locus is not known.

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The inventors herein have discovered 8 novel polymorphic sites in the NPY gene. The polymorphic sites identified by the inventors are referred to as PS1-19 to designate the order in which they are located in the gene (see Table 3 below), with the novel polymorphic sites referred to as PS1-PS5, PS9, PS13 and PS14.

5   Thus, in one embodiment, the invention provides an isolated polynucleotide comprising a polymorphic variant of the NPY gene or a fragment of the gene which contains at least one of the novel polymorphic sites described herein. The nucleotide sequence of a variant NPY gene is identical to the reference genomic sequence for those portions of the gene examined, as described in the Examples below, except that it comprises a different nucleotide at one or more of the novel

10   polymorphic sites PS1-PS5, PS9, PS13 and PS14, and may also comprise one or more additional polymorphisms selected from the group consisting of adenine at PS6, thymine at PS7, adenine at PS8, thymine at PS10, adenine at PS11, guanine at PS12, cytosine at PS15, adenine at PS16, guanine at PS17, guanine at PS18 and cytosine at PS19. Similarly, the nucleotide sequence of a variant fragment of the NPY gene is identical to the corresponding portion of the reference sequence except

15   for having a different nucleotide at one or more of the novel polymorphic sites described herein. Thus, the invention specifically does not include polynucleotides comprising a nucleotide sequence identical to the reference sequence (or other reported NPY sequences) or to portions of the reference sequence (or other reported NPY sequences), except for genotyping oligonucleotides as described below.

20   The location of a polymorphism in a variant gene or fragment is identified by aligning its sequence against SEQ ID NO:1. The polymorphism is selected from the group consisting of guanine at PS1, cytosine at PS2, thymine at PS3, adenine at PS4, thymine at PS5, cytosine at PS9, guanine at PS13 and cytosine at PS14. In a preferred embodiment, the polymorphic variant comprises a naturally-occurring isogene of the NPY gene which is defined by any one of haplotypes 1-20 shown

25   in Table 5 below.

Polymorphic variants of the invention may be prepared by isolating a clone containing the NPY gene from a human genomic library. The clone may be sequenced to determine the identity of the nucleotides at the novel polymorphic sites described herein. Any particular variant claimed herein could be prepared from this clone by performing *in vitro* mutagenesis using procedures well-

30   known in the art.

NPY isogenes may be isolated using any method that allows separation of the two "copies" of the NPY gene present in an individual, which, as readily understood by the skilled artisan, may be the same allele or different alleles. Separation methods include targeted *in vivo* cloning (TIVC) in yeast as described in WO 98/01573, U.S. Patent No. 5,866,404, and U.S. Patent No. 5,972,614.

35   Another method, which is described in U.S. Patent No. 5,972,614, uses an allele specific oligonucleotide in combination with primer extension and exonuclease degradation to generate hemizygous DNA targets. Yet other methods are single molecule dilution (SMD) as described in

Ruaño et al., Proc. Natl. Acad. Sci. 87:6296-6300, 1990; and allele specific PCR (Ruaño et al., 17 Nucleic Acids. Res. 8392, 1989; Ruaño et al., 19 Nucleic Acids Res. 6877-6882, 1991; Michalatos-Beloin et al., 24 Nucleic Acids Res. 4841-4843, 1996).

    The invention also provides NPY genome anthologies, which are collections of NPY

5   isogenes found in a given population. The population may be any group of at least two individuals, including but not limited to a reference population, a population group, a family population, a clinical population, and a same sex population. A NPY genome anthology may comprise individual NPY isogenes stored in separate containers such as microtest tubes, separate wells of a microtitre plate and the like. Alternatively, two or more groups of the NPY isogenes in the anthology may be stored in

10   separate containers. Individual isogenes or groups of isogenes in a genome anthology may be stored in any convenient and stable form, including but not limited to in buffered solutions, as DNA precipitates, freeze-dried preparations and the like. A preferred NPY genome anthology of the invention comprises a set of isogenes defined by the haplotypes shown in Table 5 below.

    An isolated polynucleotide containing a polymorphic variant nucleotide sequence of the

15   invention may be operably linked to one or more expression regulatory elements in a recombinant expression vector capable of being propagated and expressing the encoded NPY protein in a prokaryotic or a eukaryotic host cell. Examples of expression regulatory elements which may be used include, but are not limited to, the lac system, operator and promoter regions of phage lambda, yeast promoters, and promoters derived from vaccinia virus, adenovirus, retroviruses, or SV40.

20   Other regulatory elements include, but are not limited to, appropriate leader sequences, termination codons, polyadenylation signals, and other sequences required for the appropriate transcription and subsequent translation of the nucleic acid sequence in a given host cell. Of course, the correct combinations of expression regulatory elements will depend on the host system used. In addition, it is understood that the expression vector contains any additional elements necessary for its transfer to

25   and subsequent replication in the host cell. Examples of such elements include, but are not limited to, origins of replication and selectable markers. Such expression vectors are commercially available or are readily constructed using methods known to those in the art (e.g., F. Ausubel et al., 1987, in "Current Protocols in Molecular Biology", John Wiley and Sons, New York, New York). Host cells which may be used to express the variant NPY sequences of the invention include, but are not limited

30   to, eukaryotic and mammalian cells, such as animal, plant, insect and yeast cells, and prokaryotic cells, such as E. coli, or algal cells as known in the art. The recombinant expression vector may be introduced into the host cell using any method known to those in the art including, but not limited to, microinjection, electroporation, particle bombardment, transduction, and transfection using DEAE-dextran, lipofection, or calcium phosphate (see e.g., Sambrook et al. (1989) in "Molecular Cloning.

35   A Laboratory Manual", Cold Spring Harbor Press, Plainview, New York). In a preferred aspect, eukaryotic expression vectors that function in eukaryotic cells, and preferably mammalian cells, are used. Non-limiting examples of such vectors include vaccinia virus vectors, adenovirus vectors,

herpes virus vectors, and baculovirus transfer vectors. Preferred eukaryotic cell lines include COS cells, CHO cells, HeLa cells, NIH/3T3 cells, and embryonic stem cells (Thomson, J. A. et al., 1998 Science 282:1145-1147). Particularly preferred host cells are mammalian cells.

5   Polymorphic variants of fragments according to the invention comprise at least one novel polymorphism identified herein and have a length of at least 10 nucleotides and may range up to the full length of the gene. Preferably, such fragments are between 100 and 3000 nucleotides in length, and more preferably between 200 and 2000 nucleotides in length, and most preferably between 500 and 1000 nucleotides in length.

      In describing the polymorphic sites identified herein, reference is made to the sense strand of
10   the gene for convenience. However, as recognized by the skilled artisan, nucleic acid molecules containing the NPY gene may be complementary double stranded molecules and thus reference to a particular site on the sense strand refers as well to the corresponding site on the complementary antisense strand. Thus, reference may be made to the same polymorphic site on either strand and an oligonucleotide may be designed to hybridize specifically to either strand at a target region
15   containing the polymorphic site. Thus, the invention also includes single-stranded polynucleotides which are complementary to the sense strand of the NPY genomic variants described herein.

      Polynucleotides comprising a polymorphic gene variant or fragment may be useful for therapeutic purposes. For example, where a patient could benefit from expression, or increased expression, of a particular NPY protein isoform, an expression vector encoding the isoform may be
20   administered to the patient. The patient may be one who lacks the NPY isogene encoding that isoform or may already have at least one copy of that isogene.

      In other situations, it may be desirable to decrease or block expression of a particular NPY isogene. Expression of a NPY isogene may be turned off by transforming a targeted organ, tissue or cell population with an expression vector that expresses high levels of untranslatable mRNA for the
25   isogene. Alternatively, oligonucleotides directed against the regulatory regions (e.g., promoter, introns, enhancers, 3′ untranslated region) of the isogene may block transcription. Oligonucleotides targeting the transcription initiation site, e.g., between positions −10 and +10 from the start site are preferred. Similarly, inhibition of transcription can be achieved using oligonucleotides that base-pair with region(s) of the isogene DNA to form triplex DNA (see e.g., Gee et al. in Huber, B.E. and B.I.
30   Carr, Molecular and Immunologic Approaches, Futura Publishing Co., Mt. Kisco, N.Y., 1994). Antisense oligonucleotides may also be designed to block translation of NPY mRNA transcribed from a particular isogene. It is also contemplated that ribozymes may be designed that can catalyze the specific cleavage of NPY mRNA transcribed from a particular isogene.

      The oligonucleotides may be delivered to a target cell or tissue by expression from a vector
35   introduced into the cell or tissue in vivo or ex vivo. Alternatively, the oligonucleotides may be formulated as a pharmaceutical composition for administration to the patient. Oligoribonucleotides and/or oligodeoxynucleotides intended for use as antisense oligonucleotides may be modified to

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increase stability and half-life. Possible modifications include, but are not limited to phosphorothioate or 2′ O-methyl linkages, and the inclusion of nontraditional bases such as inosine and queosine, as well as acetyl-, methyl-, thio-, and similarly modified forms of adenine, cytosine, guanine, thymine, and uracil which are not as easily recognized by endogenous nucleases.

5   Effect(s) of the polymorphisms identified herein on expression of NPY may be investigated by preparing recombinant cells and/or organisms, preferably recombinant animals, containing a polymorphic variant of the NPY gene. As used herein, "expression" includes but is not limited to one or more of the following: transcription of the gene into precursor mRNA; splicing and other processing of the precursor mRNA to produce mature mRNA; mRNA stability; translation of the

10   mature mRNA into NPY protein (including codon usage and tRNA availability); and glycosylation and/or other modifications of the translation product, if required for proper expression and function.

         To prepare a recombinant cell of the invention, the desired NPY isogene may be introduced into the cell in a vector such that the isogene remains extrachromosomal. In such a situation, the gene will be expressed by the cell from the extrachromosomal location. In a preferred embodiment,

15   the NPY isogene is introduced into a cell in such a way that it recombines with the endogenous NPY gene present in the cell. Such recombination requires the occurrence of a double recombination event, thereby resulting in the desired NPY gene polymorphism. Vectors for the introduction of genes both for recombination and for extrachromosomal maintenance are known in the art, and any suitable vector or vector construct may be used in the invention. Methods such as electroporation,

20   particle bombardment, calcium phosphate co-precipitation and viral transduction for introducing DNA into cells are known in the art; therefore, the choice of method may lie with the competence and preference of the skilled practitioner. Examples of cells into which the NPY isogene may be introduced include, but are not limited to, continuous culture cells, such as COS, NIH/3T3, and primary or culture cells of the relevant tissue type, i.e., they express the NPY isogene. Such

25   recombinant cells can be used to compare the biological activities of the different protein variants.

         Recombinant organisms, i.e., transgenic animals, expressing a variant NPY gene are prepared using standard procedures known in the art. Preferably, a construct comprising the variant gene is introduced into a nonhuman animal or an ancestor of the animal at an embryonic stage, i.e., the one-cell stage, or generally not later than about the eight-cell stage. Transgenic animals carrying

30   the constructs of the invention can be made by several methods known to those having skill in the art. One method involves transfecting into the embryo a retrovirus constructed to contain one or more insulator elements, a gene or genes of interest, and other components known to those skilled in the art to provide a complete shuttle vector harboring the insulated gene(s) as a transgene, see e.g., U.S. Patent No. 5,610,053. Another method involves directly injecting a transgene into the embryo. A

35   third method involves the use of embryonic stem cells. Examples of animals into which the NPY isogenes may be introduced include, but are not limited to, mice, rats, other rodents, and nonhuman primates (see "The Introduction of Foreign Genes into Mice" and the cited references therein, In:

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Recombinant DNA, Eds. J.D. Watson, M. Gilman, J. Witkowski, and M. Zoller; W.H. Freeman and Company, New York, pages 254-272). Transgenic animals stably expressing a human NPY isogene and producing human NPY protein can be used as biological models for studying diseases related to abnormal NPY expression and/or activity, and for screening and assaying various candidate drugs,

5   compounds, and treatment regimens to reduce the symptoms or effects of these diseases.

An additional embodiment of the invention relates to pharmaceutical compositions for treating disorders affected by expression or function of a novel NPY isogene described herein. The pharmaceutical composition may comprise any of the following active ingredients: a polynucleotide comprising one of these novel NPY isogenes; an antisense oligonucleotide directed against one of the

10   novel NPY isogenes, a polynucleotide encoding such an antisense oligonucleotide, or another compound which inhibits expression of a novel NPY isogene described herein. Preferably, the composition contains the active ingredient in a therapeutically effective amount. By therapeutically effective amount is meant that one or more of the symptoms relating to disorders affected by expression or function of a novel NPY isogene is reduced and/or eliminated. The composition also

15   comprises a pharmaceutically acceptable carrier, examples of which include, but are not limited to, saline, buffered saline, dextrose, and water. Those skilled in the art may employ a formulation most suitable for the active ingredient, whether it is a polynucleotide, oligonucleotide, protein, peptide or small molecule antagonist. The pharmaceutical composition may be administered alone or in combination with at least one other agent, such as a stabilizing compound. Administration of the

20   pharmaceutical composition may be by any number of routes including, but not limited to oral, intravenous, intramuscular, intra-arterial, intramedullary, intrathecal, intraventricular, intradermal, transdermal, subcutaneous, intraperitoneal, intranasal, enteral, topical, sublingual, or rectal. Further details on techniques for formulation and administration may be found in the latest edition of Remington's Pharmaceutical Sciences (Maack Publishing Co., Easton, PA).

25   For any composition, determination of the therapeutically effective dose of active ingredient and/or the appropriate route of administration is well within the capability of those skilled in the art. For example, the dose can be estimated initially either in cell culture assays or in animal models. The animal model may also be used to determine the appropriate concentration range and route of administration. Such information can then be used to determine useful doses and routes for

30   administration in humans. The exact dosage will be determined by the practitioner, in light of factors relating to the patient requiring treatment, including but not limited to severity of the disease state, general health, age, weight and gender of the patient, diet, time and frequency of administration, other drugs being taken by the patient, and tolerance/response to the treatment.

Information on the identity of genotypes and haplotypes for the NPY gene of any particular

35   individual as well as the frequency of such genotypes and haplotypes in any particular population of individuals is expected to be useful for a variety of basic research and clinical applications. Thus, the invention also provides compositions and methods for detecting the novel NPY polymorphisms

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identified herein.

The compositions comprise at least one NPY genotyping oligonucleotide. In one

embodiment, a NPY genotyping oligonucleotide is a probe or primer capable of hybridizing to a

target region that is located close to, or that contains, one of the novel polymorphic sites described

5   herein. As used herein, the term "oligonucleotide" refers to a polynucleotide molecule having less

than about 100 nucleotides. A preferred oligonucleotide of the invention is 10 to 35 nucleotides long.

More preferably, the oligonucleotide is between 15 and 30, and most preferably, between 20 and 25

nucleotides in length. The oligonucleotide may be comprised of any phosphorylation state of

ribonucleotides, deoxyribonucleotides, and acyclic nucleotide derivatives, and other functionally

10   equivalent derivatives. Alternatively, oligonucleotides may have a phosphate-free backbone, which

may be comprised of linkages such as carboxymethyl, acetamidate, carbamate, polyamide (peptide

nucleic acid (PNA)) and the like (Varma, R. in Molecular Biology and Biotechnology, A

Comprehensive Desk Reference, Ed. R. Meyers, VCH Publishers, Inc. (1995), pages 617-620).

Oligonucleotides of the invention may be prepared by chemical synthesis using any suitable

15   methodology known in the art, or may be derived from a biological sample, for example, by

restriction digestion. The oligonucleotides may be labeled, according to any technique known in the

art, including use of radiolabels, fluorescent labels, enzymatic labels, proteins, haptens, antibodies,

sequence tags and the like.

Genotyping oligonucleotides of the invention must be capable of specifically hybridizing to a

20   target region of a NPY polynucleotide, i.e., a NPY isogene. As used herein, specific hybridization

means the oligonucleotide forms an anti-parallel double-stranded structure with the target region

under certain hybridizing conditions, while failing to form such a structure when incubated with a

non-target region or a non-NPY polynucleotide under the same hybridizing conditions. Preferably,

the oligonucleotide specifically hybridizes to the target region under conventional high stringency

25   conditions. The skilled artisan can readily design and test oligonucleotide probes and primers

suitable for detecting polymorphisms in the NPY gene using the polymorphism information provided

herein in conjunction with the known sequence information for the NPY gene and routine techniques.

A nucleic acid molecule such as an oligonucleotide or polynucleotide is said to be a "perfect"

or "complete" complement of another nucleic acid molecule if every nucleotide of one of the

30   molecules is complementary to the nucleotide at the corresponding position of the other molecule. A

nucleic acid molecule is "substantially complementary" to another molecule if it hybridizes to that

molecule with sufficient stability to remain in a duplex form under conventional low-stringency

conditions. Conventional hybridization conditions are described, for example, by Sambrook J. et al.,

in Molecular Cloning, A Laboratory Manual, 2nd Edition, Cold Spring Harbor Press, Cold Spring

35   Harbor, NY (1989) and by Haymes, B.D. et al. in Nucleic Acid Hybridization, A Practical Approach,

IRL Press, Washington, D.C. (1985). While perfectly complementary oligonucleotides are preferred

for detecting polymorphisms, departures from complete complementarity are contemplated where

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such departures do not prevent the molecule from specifically hybridizing to the target region. For
example, an oligonucleotide primer may have a non-complementary fragment at its 5′ end, with the
remainder of the primer being complementary to the target region. Alternatively, non-
complementary nucleotides may be interspersed into the oligonucleotide probe or primer as long as
5   the resulting probe or primer is still capable of specifically hybridizing to the target region.

Preferred genotyping oligonucleotides of the invention are allele-specific oligonucleotides.
As used herein, the term allele-specific oligonucleotide (ASO) means an oligonucleotide that is able,
under sufficiently stringent conditions, to hybridize specifically to one allele of a gene, or other locus,
at a target region containing a polymorphic site while not hybridizing to the corresponding region in
10   another allele(s). As understood by the skilled artisan, allele-specificity will depend upon a variety
of readily optimized stringency conditions, including salt and formamide concentrations, as well as
temperatures for both the hybridization and washing steps. Examples of hybridization and washing
conditions typically used for ASO probes are found in Kogan et al., "Genetic Prediction of
Hemophilia A" in PCR Protocols, A Guide to Methods and Applications, Academic Press, 1990 and
15   Ruano et al., 87 Proc. Natl. Acad. Sci. USA 6296-6300, 1990. Typically, an allele-specific
oligonucleotide will be perfectly complementary to one allele while containing a single mismatch for
another allele.

Allele-specific oligonucleotide probes which usually provide good discrimination between
different alleles are those in which a central position of the oligonucleotide probe aligns with the
20   polymorphic site in the target region (e.g., approximately the 7th or 8th position in a 15 mer, the 8th or
9th position in a 16mer, the 10th or 11th position in a 20 mer). A preferred ASO probe for detecting
NPY gene polymorphisms comprises a nucleotide sequence, listed 5′ to 3′, selected from the group
consisting of:

```
     TATTCATTCAACAGG   (SEQ ID NO:4)  and its complement,
25   TATTCATGCAACAGG   (SEQ ID NO:5)  and its complement,
     CAGTGGGTTGGTAGT   (SEQ ID NO:6)  and its complement,
     CAGTGGGCTGGTAGT   (SEQ ID NO:7)  and its complement,
     CGGGAGGGTTGGGGT   (SEQ ID NO:8)  and its complement,
     CGGGAGGTTTGGGGT   (SEQ ID NO:9)  and its complement,
30   CAGCCCCCTCCCCCC   (SEQ ID NO:10) and its complement,
     CAGCCCCATCCCCCC   (SEQ ID NO:11) and its complement,
     CTGTGGAGCCCTGGG   (SEQ ID NO:12) and its complement,
     CTGTGGATCCCTGGG   (SEQ ID NO:13) and its complement,
     AGAAAAGTGACCCAG   (SEQ ID NO:14) and its complement,
35   AGAAAAGCGACCCAG   (SEQ ID NO:15) and its complement,
     CGCCCGGAGCCCGCA   (SEQ ID NO:16) and its complement,
     CGCCCGGGGCCCGCA   (SEQ ID NO:17) and its complement,
     AATCCCCAAGCCCGT   (SEQ ID NO:18) and its complement, and
     AATCCCCCAGCCCGT   (SEQ ID NO:19) and its complement:
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An allele-specific oligonucleotide primer of the invention has a 3′ terminal nucleotide, or

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preferably a 3′ penultimate nucleotide, that is complementary to only one nucleotide of a particular SNP, thereby acting as a primer for polymerase-mediated extension only if the allele containing that nucleotide is present. Allele-specific oligonucleotide primers hybridizing to either the coding or noncoding strand are contemplated by the invention. A preferred ASO primer for detecting NPY gene polymorphisms comprises a nucleotide sequence, listed 5′ to 3′, selected from the group consisting of:

```
CTTGCATATTCATTC  (SEQ ID NO:20);   GTTAAACCTGTTGAA  (SEQ ID NO:21);
CTTGCATATTCATGC  (SEQ ID NO:22);   GTTAAACCTGTTGCA  (SEQ ID NO:23);
GGCACCCAGTGGGTT  (SEQ ID NO:24);   AACAGGACTACCAAC  (SEQ ID NO:25);
GGCACCCAGTGGGCT  (SEQ ID NO:26);   AACAGGACTACCAGC  (SEQ ID NO:27);
CTGGGGCGGGAGGGT  (SEQ ID NO:28);   ACCCACACCCCAACC  (SEQ ID NO:29);
CTGGGGCGGGAGGTT  (SEQ ID NO:30);   ACCCACACCCCAAAC  (SEQ ID NO:31);
CGGTTCCAGCCCCCT  (SEQ ID NO:32);   AGTGGCGGGGGGAGG  (SEQ ID NO:33);
CGGTTCCAGCCCCAT  (SEQ ID NO:34);   AGTGGCGGGGGGATG  (SEQ ID NO:35);
CCCGCTCTGTGGAGC  (SEQ ID NO:36);   CAGGGCCCCAGGGCT  (SEQ ID NO:37);
CCCGCTCTGTGGATC  (SEQ ID NO:38);   CAGGGCCCCAGGGAT  (SEQ ID NO:39);
GAAGGGAGAAAAGTG  (SEQ ID NO:40);   TTCCTGCTGGGTCAC  (SEQ ID NO:41);
GAAGGGAGAAAAGCG  (SEQ ID NO:42);   TTCCTGCTGGGTCGC  (SEQ ID NO:43);
GGACGGCGCCCGGAG  (SEQ ID NO:44);   CCACCTTGCGGGCTC  (SEQ ID NO:45);
GGACGGCGCCCGGGG  (SEQ ID NO:46);   CCACCTTGCGGGCCC  (SEQ ID NO:47);
GCTCTGAATCCCCAA  (SEQ ID NO:48);   CAACGGACGGGCTTG  (SEQ ID NO:49);
GCTCTGAATCCCCCA  (SEQ ID NO:50);   and CAACGGACGGGCTGG  (SEQ ID NO:51
).
```

Other genotyping oligonucleotides of the invention hybridize to a target region located one to several nucleotides downstream of one of the novel polymorphic sites identified herein. Such oligonucleotides are useful in polymerase-mediated primer extension methods for detecting one of the novel polymorphisms described herein and therefore such genotyping oligonucleotides are referred to herein as "primer-extension oligonucleotides". In a preferred embodiment, the 3′-terminus of a primer-extension oligonucleotide is a deoxynucleotide complementary to the nucleotide located immediately adjacent to the polymorphic site. A particularly preferred oligonucleotide primer for detecting NPY gene polymorphisms by primer extension terminates in a nucleotide sequence, listed 5′ to 3′, selected from the group consisting of:

```
GCATATTCAT  (SEQ ID NO:52);   AAACCTGTTG  (SEQ ID NO:53);
ACCCAGTGGG  (SEQ ID NO:54);   AGGACTACCA  (SEQ ID NO:55);
GGGCGGGAGG  (SEQ ID NO:56);   CACACCCCAA  (SEQ ID NO:57);
TTCCAGCCCC  (SEQ ID NO:58);   GGCGGGGGGA  (SEQ ID NO:59);
GCTCTGTGGA  (SEQ ID NO:60);   GGCCCCAGGG  (SEQ ID NO:61);
GGGAGAAAAG  (SEQ ID NO:62);   CTGCTGGGTC  (SEQ ID NO:63);
CGGCGCCCGG  (SEQ ID NO:64);   CCTTGCGGGC  (SEQ ID NO:65);
CTGAATCCCC  (SEQ ID NO:66);   and CGGACGGGCT  (SEQ ID NO:67).
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In some embodiments, a composition contains two or more differently labeled genotyping oligonucleotides for simultaneously probing the identity of nucleotides at two or more polymorphic

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sites. It is also contemplated that primer compositions may contain two or more sets of allele-specific primer pairs to allow simultaneous targeting and amplification of two or more regions containing a polymorphic site.

NPY genotyping oligonucleotides of the invention may also be immobilized on or
5   synthesized on a solid surface such as a microchip, bead, or glass slide (see, e.g., WO 98/20020 and WO 98/20019). Such immobilized genotyping oligonucleotides may be used in a variety of polymorphism detection assays, including but not limited to probe hybridization and polymerase extension assays. Immobilized NPY genotyping oligonucleotides of the invention may comprise an ordered array of oligonucleotides designed to rapidly screen a DNA sample for polymorphisms in
10   multiple genes at the same time.

In another embodiment, the invention provides a kit comprising at least two genotyping oligonucleotides packaged in separate containers. The kit may also contain other components such as hybridization buffer (where the oligonucleotides are to be used as a probe) packaged in a separate container. Alternatively, where the oligonucleotides are to be used to amplify a target region, the kit
15   may contain, packaged in separate containers, a polymerase and a reaction buffer optimized for primer extension mediated by the polymerase, such as PCR.

The above described oligonucleotide compositions and kits are useful in methods for genotyping and/or haplotyping the NPY gene in an individual. As used herein, the terms "NPY genotype" and "NPY haplotype" mean the genotype or haplotype contains the nucleotide pair or
20   nucleotide, respectively, that is present at one or more of the novel polymorphic sites described herein and may optionally also include the nucleotide pair or nucleotide present at one or more additional polymorphic sites in the NPY gene. The additional polymorphic sites may be currently known polymorphic sites or sites that are subsequently discovered.

One embodiment of the genotyping method involves isolating from the individual a nucleic
25   acid mixture comprising the two copies of the NPY gene, or a fragment thereof, that are present in the individual, and determining the identity of the nucleotide pair at one or more of the polymorphic sites selected from PS1-PS5, PS9, PS13 and PS14 in the two copies to assign a NPY genotype to the individual. As will be readily understood by the skilled artisan, the two "copies" of a gene in an individual may be the same allele or may be different alleles. In a preferred embodiment of the
30   genotyping method, the identity of the nucleotide pair at one or more of the polymorphic sites selected from the group consisting of PS6-PS8, PS10-PS12 and PS15-PS19 is also determined. In a particularly preferred embodiment, the genotyping method comprises determining the identity of the nucleotide pair at each of PS1-19.

Typically, the nucleic acid mixture is isolated from a biological sample taken from the
35   individual, such as a blood sample or tissue sample. Suitable tissue samples include whole blood, semen, saliva, tears, urine, fecal material, sweat, buccal, skin and hair. The nucleic acid mixture may be comprised of genomic DNA, mRNA, or cDNA and, in the latter two cases, the biological sample

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must be obtained from an organ in which the NPY gene is expressed. Furthermore it will be understood by the skilled artisan that mRNA or cDNA preparations would not be used to detect polymorphisms located in introns or in 5′ and 3′ nontranscribed regions. If a NPY gene fragment is isolated, it must contain the polymorphic site(s) to be genotyped.

5   One embodiment of the haplotyping method comprises isolating from the individual a nucleic acid molecule containing only one of the two copies of the NPY gene, or a fragment thereof, that is present in the individual and determining in that copy the identity of the nucleotide at one or more of the polymorphic sites PS1-PS5, PS9, PS13 and PS14 in that copy to assign a NPY haplotype to the individual. The nucleic acid may be isolated using any method capable of separating the two

10   copies of the NPY gene or fragment such as one of the methods described above for preparing NPY isogenes, with targeted *in vivo* cloning being the preferred approach. As will be readily appreciated by those skilled in the art, any individual clone will only provide haplotype information on one of the two NPY gene copies present in an individual. If haplotype information is desired for the individual's other copy, additional NPY clones will need to be examined. Typically, at least five

15   clones should be examined to have more than a 90% probability of haplotyping both copies of the NPY gene in an individual. In some embodiments, the haplotyping method also comprises identifying the nucleotide at one or more of the polymorphic sites PS6-PS8, PS10-PS12 and PS15-PS19. In a particularly preferred embodiment, the nucleotide at each of PS1-19 is identified.

           In a preferred embodiment, a NPY haplotype pair is determined for an individual by

20   identifying the phased sequence of nucleotides at one or more of the polymorphic sites selected from PS1-PS5, PS9, PS13 and PS14 in each copy of the NPY gene that is present in the individual. In a particularly preferred embodiment, the haplotyping method comprises identifying the phased sequence of nucleotides at each of PS1-19 in each copy of the NPY gene. When haplotyping both copies of the gene, the identifying step is preferably performed with each copy of the gene being

25   placed in separate containers. However, it is also envisioned that if the two copies are labeled with different tags, or are otherwise separately distinguishable or identifiable, it could be possible in some cases to perform the method in the same container. For example, if first and second copies of the gene are labeled with different first and second fluorescent dyes, respectively, and an allele-specific oligonucleotide labeled with yet a third different fluorescent dye is used to assay the polymorphic

30   site(s), then detecting a combination of the first and third dyes would identify the polymorphism in the first gene copy while detecting a combination of the second and third dyes would identify the polymorphism in the second gene copy.

           In both the genotyping and haplotyping methods, the identity of a nucleotide (or nucleotide pair) at a polymorphic site(s) may be determined by amplifying a target region(s) containing the

35   polymorphic site(s) directly from one or both copies of the NPY gene, or fragment thereof, and the sequence of the amplified region(s) determined by conventional methods. It will be readily appreciated by the skilled artisan that only one nucleotide will be detected at a polymorphic site in

                                              19

individuals who are homozygous at that site, while two different nucleotides will be detected if the individual is heterozygous for that site. The polymorphism may be identified directly, known as positive-type identification, or by inference, referred to as negative-type identification. For example, where a SNP is known to be guanine and cytosine in a reference population, a site may be positively

5   determined to be either guanine or cytosine for an individual homozygous at that site, or both guanine and cytosine, if the individual is heterozygous at that site. Alternatively, the site may be negatively determined to be not guanine (and thus cytosine/cytosine) or not cytosine (and thus guanine/guanine).

In addition, the identity of the allele(s) present at any of the novel polymorphic sites

10   described herein may be indirectly determined by genotyping a polymorphic site not disclosed herein that is in linkage disequilibrium with the polymorphic site that is of interest. Two sites are said to be in linkage disequilibrium if the presence of a particular variant at one site enhances the predictability of another variant at the second site (Stevens, JC 1999, *Mol. Diag.* 4: 309-17). Polymorphic sites in linkage disequilibrium with the presently disclosed polymorphic sites may be located in regions of

15   the gene or in other genomic regions not examined herein. Genotyping of a polymorphic site in linkage disequilibrium with the novel polymorphic sites described herein may be performed by, but is not limited to, any of the above-mentioned methods for detecting the identity of the allele at a polymorphic site.

The target region(s) may be amplified using any oligonucleotide-directed amplification

20   method, including but not limited to polymerase chain reaction (PCR) (U.S. Patent No. 4,965,188), ligase chain reaction (LCR) (Barany et al., Proc. Natl. Acad. Sci. USA 88:189-193, 1991; WO90/01069), and oligonucleotide ligation assay (OLA) (Landegren et al., Science 241:1077-1080, 1988). Oligonucleotides useful as primers or probes in such methods should specifically hybridize to a region of the nucleic acid that contains or is adjacent to the polymorphic site. Typically, the

25   oligonucleotides are between 10 and 35 nucleotides in length and preferably, between 15 and 30 nucleotides in length. Most preferably, the oligonucleotides are 20 to 25 nucleotides long. The exact length of the oligonucleotide will depend on many factors that are routinely considered and practiced by the skilled artisan.

Other known nucleic acid amplification procedures may be used to amplify the target region

30   including transcription-based amplification systems (U.S. Patent No. 5,130,238; EP 329,822; U.S. Patent No. 5,169,766, WO89/06700) and isothermal methods (Walker et al., *Proc. Natl. Acad. Sci. USA* 89:392-396, 1992).

A polymorphism in the target region may also be assayed before or after amplification using one of several hybridization-based methods known in the art. Typically, allele-specific

35   oligonucleotides are utilized in performing such methods. The allele-specific oligonucleotides may be used as differently labeled probe pairs, with one member of the pair showing a perfect match to one variant of a target sequence and the other member showing a perfect match to a different variant.

In some embodiments, more than one polymorphic site may be detected at once using a set of allele-specific oligonucleotides or oligonucleotide pairs. Preferably, the members of the set have melting temperatures within 5°C, and more preferably within 2°C, of each other when hybridizing to each of the polymorphic sites being detected.

5   Hybridization of an allele-specific oligonucleotide to a target polynucleotide may be performed with both entities in solution, or such hybridization may be performed when either the oligonucleotide or the target polynucleotide is covalently or noncovalently affixed to a solid support. Attachment may be mediated, for example, by antibody-antigen interactions, poly-L-Lys, streptavidin or avidin-biotin, salt bridges, hydrophobic interactions, chemical linkages, UV cross-

10   linking baking, etc. Allele-specific oligonucleotides may be synthesized directly on the solid support or attached to the solid support subsequent to synthesis. Solid-supports suitable for use in detection methods of the invention include substrates made of silicon, glass, plastic, paper and the like, which may be formed, for example, into wells (as in 96-well plates), slides, sheets, membranes, fibers, chips, dishes, and beads. The solid support may be treated, coated or derivatized to facilitate the

15   immobilization of the allele-specific oligonucleotide or target nucleic acid.

The genotype or haplotype for the NPY gene of an individual may also be determined by hybridization of a nucleic acid sample containing one or both copies of the gene to nucleic acid arrays and subarrays such as described in WO 95/11995. The arrays would contain a battery of allele-specific oligonucleotides representing each of the polymorphic sites to be included in the

20   genotype or haplotype.

The identity of polymorphisms may also be determined using a mismatch detection technique, including but not limited to the RNase protection method using riboprobes (Winter et al., Proc. Natl. Acad. Sci. USA 82:7575, 1985; Meyers et al., Science 230:1242, 1985) and proteins which recognize nucleotide mismatches, such as the E. coli mutS protein (Modrich, P. Ann. Rev.

25   Genet. 25:229-253, 1991). Alternatively, variant alleles can be identified by single strand conformation polymorphism (SSCP) analysis (Orita et al., Genomics 5:874-879, 1989; Humphries et al., in Molecular Diagnosis of Genetic Diseases, R. Elles, ed., pp. 321-340, 1996) or denaturing gradient gel electrophoresis (DGGE) (Wartell et al., Nucl. Acids Res. 18:2699-2706, 1990; Sheffield et al., Proc. Natl. Acad. Sci. USA 86:232-236, 1989).

30   A polymerase-mediated primer extension method may also be used to identify the polymorphism(s). Several such methods have been described in the patent and scientific literature and include the "Genetic Bit Analysis" method (WO92/15712) and the ligase/polymerase mediated genetic bit analysis (U.S. Patent 5,679,524. Related methods are disclosed in WO91/02087, WO90/09455, WO95/17676, U.S. Patent Nos. 5,302,509, and 5,945,283. Extended primers

35   containing a polymorphism may be detected by mass spectrometry as described in U.S. Patent No. 5,605,798. Another primer extension method is allele-specific PCR (Ruaño et al., Nucl. Acids Res. 17:8392, 1989; Ruaño et al., Nucl. Acids Res. 19, 6877-6882, 1991; WO 93/22456; Turki et al., J.

Clin. Invest. 95:1635-1641, 1995). In addition, multiple polymorphic sites may be investigated by simultaneously amplifying multiple regions of the nucleic acid using sets of allele-specific primers as described in Wallace et al. (WO89/10414).

In another aspect of the invention, an individual's NPY haplotype pair is predicted from its
5   NPY genotype using information on haplotype pairs known to exist in a reference population. In its broadest embodiment, the haplotyping prediction method comprises identifying a NPY genotype for the individual at two or more polymorphic sites selected from PS1-PS5, PS9, PS13 and PS14, enumerating all possible haplotype pairs which are consistent with the genotype, accessing data containing NPY haplotype pairs identified in a reference population, and assigning a haplotype pair
10 to the individual that is consistent with the data. In one embodiment, the reference haplotype pairs include the NPY haplotype pairs shown in Table 4.

Generally, the reference population should be composed of randomly-selected individuals representing the major ethnogeographic groups of the world. A preferred reference population for use in the methods of the present invention comprises an approximately equal number of individuals
15 from Caucasian, African American, Asian and Hispanic-Latino population groups with the minimum number of each group being chosen based on how rare a haplotype one wants to be guaranteed to see. For example, if one wants to have a q% chance of not missing a haplotype that exists in the population at a p% frequency of occurring in the reference population, the number of individuals (n) who must be sampled is given by $2n=\log(1-q)/\log(1-p)$ where p and q are expressed as fractions. A
20 preferred reference population allows the detection of any haplotype whose frequency is at least 10% with about 99% certainty and comprises about 20 unrelated individuals from each of the four population groups named above. A particularly preferred reference population includes a 3-generation family representing one or more of the four population groups to serve as controls for checking quality of haplotyping procedures.

25 In a preferred embodiment, the haplotype frequency data for each ethnogeographic group is examined to determine whether it is consistent with Hardy-Weinberg equilibrium. Hardy-Weinberg equilibrium (D.L. Hartl et al., Principles of Population Genomics, Sinauer Associates (Sunderland, MA), 3rd Ed., 1997) postulates that the frequency of finding the haplotype pair H_1 / H_2 is equal to $p_{H-W}(H_1 / H_2) = 2p(H_1)p(H_2)$ if $H_1 \neq H_2$ and $p_{H-W}(H_1 / H_2) = p(H_1)p(H_2)$ if $H_1 = H_2$.

30 A statistically significant difference between the observed and expected haplotype frequencies could be due to one or more factors including significant inbreeding in the population group, strong selective pressure on the gene, sampling bias, and/or errors in the genotyping process. If large deviations from Hardy-Weinberg equilibrium are observed in an ethnogeographic group, the number of individuals in that group can be increased to see if the deviation is due to a sampling bias. If a
35 larger sample size does not reduce the difference between observed and expected haplotype pair frequencies, then one may wish to consider haplotyping the individual using a direct haplotyping

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method such as, for example, CLASPER System™ technology (U.S. Patent No. 5,866,404), SMD, or allele-specific long-range PCR (Michalotos-Beloin et al., Nucleic Acids Res. 24:4841-4843, 1996). A preferred process for predicting NPY haplotype pairs from NPY genotypes is described in copending U.S. Provisional Application Serial No. 60/198,340.

5   In one embodiment of this method for predicting a NPY haplotype pair, the assigning step involves performing the following analysis. First, each of the possible haplotype pairs is compared to the haplotype pairs in the reference population. Generally, only one of the haplotype pairs in the reference population matches a possible haplotype pair and that pair is assigned to the individual. Occasionally, only one haplotype represented in the reference haplotype pairs is consistent with a

10   possible haplotype pair for an individual, and in such cases the individual is assigned a haplotype pair containing this known haplotype and a new haplotype derived by subtracting the known haplotype from the possible haplotype pair. In rare cases, either no haplotypes in the reference population are consistent with the possible haplotype pairs, or alternatively, multiple reference haplotype pairs are consistent with the possible haplotype pairs. In such cases, the individual is preferably haplotyped

15   using a direct molecular haplotyping method such as, for example, CLASPER System™ technology (U.S. Patent No. 5,866,404), SMD, or allele-specific long-range PCR (Michalotos-Beloin et al., Nucleic Acids Res. 24:4841-4843, 1996).

          The invention also provides a method for determining the frequency of a NPY genotype or NPY haplotype in a population. The method comprises determining the genotype or the haplotype

20   pair for the NPY gene that is present in each member of the population, wherein the genotype or haplotype comprises the nucleotide pair or nucleotide detected at one or more of the polymorphic sites PS1-PS5, PS9, PS13 and PS14 in the NPY gene; and calculating the frequency any particular genotype or haplotype is found in the population. The population may be a reference population, a family population, a same sex population, a population group, a trait population (e.g., a group of

25   individuals exhibiting a trait of interest such as a medical condition or response to a therapeutic treatment). A preferred process for predicting NPY haplotype pairs from NPY genotypes is described in pending U.S. Provisional Application Serial No. 60/198,340.

          In another aspect of the invention, frequency data for NPY genotypes and/or haplotypes found in a reference population are used in a method for identifying an association between a trait

30   and a NPY genotype or a NPY haplotype. The trait may be any detectable phenotype, including but not limited to susceptibility to a disease or response to a treatment. The method involves obtaining data on the frequency of the genotype(s) or haplotype(s) of interest in a reference population as well as in a population exhibiting the trait. Frequency data for one or both of the reference and trait populations may be obtained by genotyping or haplotyping each individual in the populations using

35   one of the methods described above. The haplotypes for the trait population may be determined directly or, alternatively, by the predictive genotype to haplotype approach described above. In another embodiment, the frequency data for the reference and/or trait populations is obtained by

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accessing previously determined frequency data, which may be in written or electronic form. For example, the frequency data may be present in a database that is accessible by a computer. Once the frequency data is obtained, the frequencies of the genotype(s) or haplotype(s) of interest in the reference and trait populations are compared. In a preferred embodiment, the frequencies of all

5   genotypes and/or haplotypes observed in the populations are compared. If a particular genotype or haplotype for the NPY gene is more frequent in the trait population than in the reference population at a statistically significant amount, then the trait is predicted to be associated with that NPY genotype or haplotype. Preferably, the NPY genotype or haplotype being compared in the trait and reference populations is selected from the full-genotypes and full-haplotypes shown in Tables 4 and

10   5, respectively, or from sub-genotypes and sub-haplotypes derived from these genotypes and haplotypes.

         In a preferred embodiment of the method, the trait of interest is a clinical response exhibited by a patient to some therapeutic treatment, for example, response to a drug targeting NPY or response to a therapeutic treatment for a medical condition. As used herein, "medical condition"

15   includes but is not limited to any condition or disease manifested as one or more physical and/or psychological symptoms for which treatment is desirable, and includes previously and newly identified diseases and other disorders. As used herein the term "clinical response" means any or all of the following: a quantitative measure of the response, no response, and adverse response (i.e., side effects).

20   In order to deduce a correlation between clinical response to a treatment and a NPY genotype or haplotype, it is necessary to obtain data on the clinical responses exhibited by a population of individuals who received the treatment, hereinafter the "clinical population". This clinical data may be obtained by analyzing the results of a clinical trial that has already been run and/or the clinical data may be obtained by designing and carrying out one or more new clinical trials. As used herein,

25   the term "clinical trial" means any research study designed to collect clinical data on responses to a particular treatment, and includes but is not limited to phase I, phase II and phase III clinical trials. Standard methods are used to define the patient population and to enroll subjects.

         It is preferred that the individuals included in the clinical population have been graded for the existence of the medical condition of interest. This is important in cases where the symptom(s) being

30   presented by the patients can be caused by more than one underlying condition, and where treatment of the underlying conditions are not the same. An example of this would be where patients experience breathing difficulties that are due to either asthma or respiratory infections. If both sets were treated with an asthma medication, there would be a spurious group of apparent non-responders that did not actually have asthma. These people would affect the ability to detect any correlation

35   between haplotype and treatment outcome. This grading of potential patients could employ a standard physical exam or one or more lab tests. Alternatively, grading of patients could use haplotyping for situations where there is a strong correlation between haplotype pair and disease

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susceptibility or severity.

The therapeutic treatment of interest is administered to each individual in the trial population and each individual's response to the treatment is measured using one or more predetermined criteria. It is contemplated that in many cases, the trial population will exhibit a range of responses and that

5 the investigator will choose the number of responder groups (e.g., low, medium, high) made up by the various responses. In addition, the NPY gene for each individual in the trial population is genotyped and/or haplotyped, which may be done before or after administering the treatment.

After both the clinical and polymorphism data have been obtained, correlations between individual response and NPY genotype or haplotype content are created. Correlations may be

10 produced in several ways. In one method, individuals are grouped by their NPY genotype or haplotype (or haplotype pair) (also referred to as a polymorphism group), and then the averages and standard deviations of clinical responses exhibited by the members of each polymorphism group are calculated.

These results are then analyzed to determine if any observed variation in clinical response

15 between polymorphism groups is statistically significant. Statistical analysis methods which may be used are described in L.D. Fisher and G. vanBelle, "Biostatistics: A Methodology for the Health Sciences", Wiley-Interscience (New York) 1993. This analysis may also include a regression calculation of which polymorphic sites in the NPY gene give the most significant contribution to the differences in phenotype. One regression model useful in the invention is described in PCT

20 Application Serial No. PCT/US00/17540, entitled "Methods for Obtaining and Using Haplotype Data".

A second method for finding correlations between NPY haplotype content and clinical responses uses predictive models based on error-minimizing optimization algorithms. One of many possible optimization algorithms is a genetic algorithm (R. Judson, "Genetic Algorithms and Their

25 Uses in Chemistry" in Reviews in Computational Chemistry, Vol. 10, pp. 1-73, K. B. Lipkowitz and D. B. Boyd, eds. (VCH Publishers, New York, 1997). Simulated annealing (Press et al., "Numerical Recipes in C: The Art of Scientific Computing", Cambridge University Press (Cambridge) 1992, Ch. 10), neural networks (E. Rich and K. Knight, "Artificial Intelligence", 2nd Edition (McGraw-Hill, New York, 1991, Ch. 18), standard gradient descent methods (Press et al., supra Ch. 10), or other

30 global or local optimization approaches (see discussion in Judson, supra) could also be used. Preferably, the correlation is found using a genetic algorithm approach as described in PCT Application Serial No. PCT/US00/17540.

Correlations may also be analyzed using analysis of variation (ANOVA) techniques to determine how much of the variation in the clinical data is explained by different subsets of the

35 polymorphic sites in the NPY gene. As described in PCT Application Serial No. PCT/US00/17540, ANOVA is used to test hypotheses about whether a response variable is caused by or correlated with one or more traits or variables that can be measured (Fisher and vanBelle, supra, Ch. 10).

25

From the analyses described above, a mathematical model may be readily constructed by the skilled artisan that predicts clinical response as a function of NPY genotype or haplotype content. Preferably, the model is validated in one or more follow-up clinical trials designed to test the model.

The identification of an association between a clinical response and a genotype or haplotype

5   (or haplotype pair) for the NPY gene may be the basis for designing a diagnostic method to determine those individuals who will or will not respond to the treatment, or alternatively, will respond at a lower level and thus may require more treatment, i.e., a greater dose of a drug. The diagnostic method may take one of several forms: for example, a direct DNA test (i.e., genotyping or haplotyping one or more of the polymorphic sites in the NPY gene), a serological test, or a physical

10   exam measurement. The only requirement is that there be a good correlation between the diagnostic test results and the underlying NPY genotype or haplotype that is in turn correlated with the clinical response. In a preferred embodiment, this diagnostic method uses the predictive haplotyping method described above.

Any or all analytical and mathematical operations involved in practicing the methods of the

15   present invention may be implemented by a computer. In addition, the computer may execute a program that generates views (or screens) displayed on a display device and with which the user can interact to view and analyze large amounts of information relating to the NPY gene and its genomic variation, including chromosome location, gene structure, and gene family, gene expression data, polymorphism data, genetic sequence data, and clinical data population data (e.g., data on

20   ethnogeographic origin, clinical responses, genotypes, and haplotypes for one or more populations). The NPY polymorphism data described herein may be stored as part of a relational database (e.g., an instance of an Oracle database or a set of ASCII flat files). These polymorphism data may be stored on the computer's hard drive or may, for example, be stored on a CD ROM or on one or more other storage devices accessible by the computer. For example, the data may be stored on one or more

25   databases in communication with the computer via a network.

Preferred embodiments of the invention are described in the following examples. Other embodiments within the scope of the claims herein will be apparent to one skilled in the art from consideration of the specification or practice of the invention as disclosed herein. It is intended that the specification, together with the examples, be considered exemplary only, with the scope and spirit

30   of the invention being indicated by the claims which follow the examples.

EXAMPLES

The Examples herein are meant to exemplify the various aspects of carrying out the invention and are not intended to limit the scope of the invention in any way. The Examples do not

35   include detailed descriptions for conventional methods employed, such as in the performance of genomic DNA isolation, PCR and sequencing procedures. Such methods are well-known to those skilled in the art and are described in numerous publications, for example, Sambrook, Fritsch, and

Maniatis, "Molecular Cloning: A Laboratory Manual", 2[nd] Edition, Cold Spring Harbor Laboratory

Press, USA, (1989).

EXAMPLE 1

5   This example illustrates examination of various regions of the NPY gene for polymorphic

sites.

Amplification of Target Regions

10   The following target regions of the NPY gene were amplified using the PCR primer pairs

listed below, with the sequences presented in the 5′ to 3′ direction and nucleotide positions shown

for each region corresponding to the reverse complement of the indicated GenBank Accession No.

Accession No. ACC004485.1
Fragment 1
15 Forward primer:
154135-154158   5'- GGAGCATTCATTCACGGAACTTTC -3' (SEQ ID NO:68)
Reverse primer:
Complement of 154540-154520 5'- GGCTCGGTCGAGACATGAGTG -3' (SEQ ID NO:69)
PCR Product   406 nt

20
Fragment 2
Forward primer:
154364-154387   5'- CATTTAACCACCGTCACTTTGGAC -3' (SEQ ID NO:70)
Reverse primer:
25 Complement of 154901-154879 5'- AGTCCTGCCAGAGATAGGAGCAG -3' (SEQ ID NO:71)
PCR Product   538 nt

Fragment 3
Forward primer:
30 154726-154747   5'- CCAGTTGCCTCACTCCAACAGC -3' (SEQ ID NO:72)
Reverse primer:
Complement of 155161-155142 5'- TGACTCCCACCCGCCACTTC -3' (SEQ ID NO:73)
PCR Product   436 nt

35 Fragment 4
Forward primer:
154967-154987   5'- ACTTGGGCTTTGGTGTTGTCC -3' (SEQ ID NO:74)
Reverse primer:
Complement of 155532-155511 5'- AATCAGAGGGGAGCAGAGGAAC -3' (SEQ ID NO:75)
40 PCR Product   566 nt

Fragment 5
Forward primer:
155377-155399   5'- GAACTTGCTAGAGACGCAGCCTC -3' (SEQ ID NO:76)
45 Reverse primer:
Complement of 155881-155861 5'- CTTCAGGGGATCAACGCTGAC -3' (SEQ ID NO:77)
PCR Product   505 nt

Fragment 6
Forward primer:
155760-155780   5'- TGGATTCTTGGGCTCCAAATC -3' (SEQ ID NO:78)
Reverse primer:
5   Complement of 156262-156243 5'- GCTTGGGGATTCAGAGCACC -3' (SEQ ID NO:79)
PCR Product   503 nt

Fragment 7
Forward primer:
10   156031-156053   5'- GCTTCTTGGTCCCTGAGACTTCG -3' (SEQ ID NO:80)
Reverse primer:
Complement of 156556-156536 5'- CCCTTTCCCTAACATCCCCAG -3' (SEQ ID NO:81)
PCR Product   526 nt

15   Fragment 8
Forward primer:
160417-160442   5'- GCTTGCTTGTTACAGATGAACACCTG -3' (SEQ ID NO:82)
Reverse primer:
Complement of 160870-160848 5'- CGTCAAAGGAGCACCAAATAACC -3' (SEQ ID NO:83)
20   PCR Product   454 nt

Fragment 9
Forward primer:
162593-162616   5'- CAGGAAACTTTTCAACAGTTCCCG -3' (SEQ ID NO:84)
25   Reverse primer:
Complement of 163029-163009 5'- GCGAAACGAACCCTGAATCTG -3' (SEQ ID NO:85)
PCR Product   437 nt

30   These primer pairs were used in PCR reactions containing genomic DNA isolated from

immortalized cell lines for each member of the Index Repository. The PCR reactions were carried

out under the following conditions:

Reaction volume	= 20 μl
10 x Advantage 2 Polymerase reaction buffer (Clontech)	= 2 μl
35 100 ng of human genomic DNA	= 1 μl
10 mM dNTP	= 0.4 μl
Advantage 2 Polymerase enzyme mix (Clontech)	= 0.2 μl
Forward Primer (10 μM)	= 0.4 μl
Reverse Primer (10 μM)	= 0.4 μl
40 Water	=15.6μl

Amplification profile:
94°C - 2 min.   1 cycle

45   94°C - 30 sec.
70°C - 45 sec.   } 10 cycles
72°C - 1 min.

50   94°C - 30 sec.
64°C - 45 sec.   } 35 cycles
72°C - 1 min.

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Sequencing of PCR Products

The PCR products were purified by Solid Phase Reversible Immobilization using the protocol developed by the Whitehead Genome Center. A detailed protocol can be found at http://www.genome.wi.mit.edu/sequencing/protocols/pure/SPRI_pcr.html.

5   Briefly, five µl of carboxyl coated magnetic beads (10 mg/ml) and 60 µl of HYB BUFFER (2.5M NaCl/20% PEG 8000) were added to each PCR reaction mixture (20 µl). The reaction mixture was mixed well and incubated at room temperature (RT) for 10 min. The microtitre plate was placed on a magnet for 2 min and the beads washed twice with 150 µl of 70% EtOH. The beads were air dried for 2 min and the DNA was eluted in 25 µl of distilled water and incubated at RT for 5

10   min. The beads were magnetically separated and the supernatant removed for testing and sequencing.

The purified PCR products were sequenced in both directions using the primer sets described previously or those listed below, in the 5′ to 3′ direction.

Fragment 7
15   Reverse sequencing primer:
Complement of 156535-156515 5′- GATCTCCTGGTGTGCAGGCAC -3′ (SEQ ID NO:86)

Fragment 9
Forward sequencing primer
20   162638-162661   5′- AAATGTCTCACCCTTGCTCATACC -3′ (SEQ ID NO:87)
Reverse sequencing primer
Complement of 163023-163000 5′- CGAACCCTGAATCTGCTAGTTGAG -3′ (SEQ ID NO:88)

25   ## Analysis of Sequences for Polymorphic Sites

Sequences were analyzed for the presence of polymorphisms using the Polyphred program (Nickerson et al., Nucleic Acids Res. 14:2745-2751, 1997). The presence of a polymorphism was confirmed on both strands. The polymorphisms and their locations in the NPY gene are listed in Table 3 below.

Table 3.   Polymorphic Sites Identified in the NPY Gene

	Polymorphic Site Number	PolyId	Nucleotide Position in GenBank	Nucleotide Position in Fig 1	Reference. Allele	Variant Allele
5	PS1	98214	31293(Acc#AC004485.1)	154224	T	G
	PS2	98218	30676(Acc#AC004485.1)	154841	T	C
	PS3	98219	30454(Acc#AC004485.1)	155063	G	T
	PS4	98220	30398(Acc#AC004485.1)	155119	C	A
	PS5	98222	30105(Acc#AC004485.1)	155412	G	T
10	PS6 [R]	98223	30104(Acc#AC004485.1)	155413	C	A
	PS7 [R]	98224	30097(Acc#AC004485.1)	155420	G	T
	PS8 [R]	98227	29955(Acc#AC004485.1)	155562	G	A
	PS9	98230	29563(Acc#AC004485.1)	155954	T	C
	PS10 [R]	98231	29498(Acc#AC004485.1)	156019	C	T
15	PS11 [R]	98232	29478(Acc#AC004485.1)	156039	G	A
	PS12 [R]	98233	29416(Acc#AC004485.1)	156101	A	G
	PS13	98234	29400(Acc#AC004485.1)	156117	A	G
	PS14	98235	29258(Acc#AC004485.1)	156259	A	C
	PS15 [R]	98236	29207(Acc#AC004485.1)	156310	T	C
20	PS16 [R]	98237	29143(Acc#AC004485.1)	156374	G	A
	PS17 [R]	98238	22735(Acc#AC004485.1)	162782	A	G
	PS18 [R]	98239	22600(Acc#AC004485.1)	162917	A	G
	PS19 [R]	98240	22569(Acc#AC004485.1)	162948	T	C

25   [R] Reported previously in literature

EXAMPLE 2

This example illustrates analysis of the NPY polymorphisms identified in the Index

Repository for human genotypes and haplotypes.

30   The different genotypes containing these polymorphisms that were observed in the reference

population are shown in Table 4 below, with the haplotype pair indicating the combination of

haplotypes determined for the individual using the haplotype derivation protocol described below. In

Table 4, homozygous positions are indicated by one nucleotide and heterozygous positions are

indicated by two nucleotides. Missing nucleotides in any given genotype in Table 4 were inferred

35   based on linkage disequilibrium and/or Mendelian inheritance.

Table 4(Part1). Genotypes and Haplotype Pairs Observed for NPY Gene Polymorphic Sites

Genotype Number Pair	PS1	PS2	PS3	PS4	PS5	PS6	PS7	PS8	PS9	PS10	HAP	
1	T	T	G	C	G	C	G	G	T	C	9	9
2	T	C	G	C	G	C	T	G	T	C	4	4
3	T	T	G	C	G	C	G	G	T	T	13	13
4	T	C/T	G	C	G	C	T/G	G	T	C/T	4	14
5	T	C	G	C	G	C	T	G	T	C	4	6
6	T	T	G	C	G	C	G	G	T	T	13	14
7	T	C/T	G	C	G	C	T/G	G	T	C/T	4	17
8	T	C/T	G	C	G/T	C	T/G	G	T	C	4	19
9	T	C/T	G	C	G	C	T/G	G	T	C	4	11
10	T	T	G	C	G	C	G	G	T	T	13	16
11	T	C/T	G	C	G	C	T/G	G	T	C/T	4	15
12	T	T	G	C	G	C/A	G	G	T	T	13	7
13	T	C/T	G	C	G	C	T	G	T/C	C	4	18
14	T	C	G	C	G	C	T	G/A	T	C	4	3
15	T	C	G	C	G	C	T	G	T	C	4	5
16	T	T	G	C	G	C	G/T	G	T/C	T/C	13	18
17	T	C/T	G/T	C	G	C	T/G	G	T	C/T	4	20
18	T	C	G	C/A	G	C	T	G	T	C	4	2
19	T	C/T	G	C	G	C	T/G	G	T	C/T	4	12
20	T	T	G	C	G	C	G	G	T	T/C	13	9
21	T	C/T	G	C	G	C	T/G	G	T	C	4	9
22	T	T	G	C	G	C	G	G	T	T/C	13	8
23	T	T	G	C	G	C	G	G	T	T/C	13	10
24	T	C/T	G	C	G	C	T/G	G	T	C	4	8
25	T	T	G	C	G	C	G	G	T	C	9	8
26	T	C/T	G	C	G	C	T/G	G	T	C/T	4	13
27	T/G	T/C	G	C	G	C	G	G	T	C/T	9	1

Table 4(Part2). Genotypes and Haplotype Pairs Observed for NPY Gene

Genotype Number	PS11	PS12	PS13	PS14	PS15	PS16	PS17	PS18	PS19	HAP	Pair
1	G	A	A	A	T	G	A	A	T	9	9
2	G	A	A	A	T	G	A	A	T	4	4
3	A	G	A	A	T	G	A	A	T	13	13
4	G/A	A/G	A	A	T	G	A	A/G	T	4	14
5	G	A	A/G	A	T	G	A	A	T	4	6
6	A	G	A	A	T	G	A	A/G	T	13	14
7	G	A/G	A	A	T	G/A	A/G	A	T	4	17
8	G/A	A/G	A	A	T	G	A	A	T/C	4	19
9	G	A	A	A	T	G	A/G	A	T	4	11
10	A/G	G	A	A	T	G/A	A	A	T/C	13	16
11	G/A	A/G	A	A	T	G	A/G	A	T	4	15
12	A	G.	A	A	T	G	A	A	T	13	7
13	G	A	A	A	T	G	A	A	T	4	18
14	G	A	A	A	T	G	A	A	T	4	3
15	G	A	A	A/C	T	G	A	A	T	4	5
16	A/G	G/A	A	A	T	G	A	A	T	13	18
17	G/A	A/G	A	A	T	G	A	A	T	4	20
18	G	A	A	A	T	G	A	A	T	4	2
19	G/A	A/G	A	A	T	G	A	A	T/C	4	12
20	A/G	G/A	A	A	T	G	A	A.	T	13	9
21	G	A	A	A	T	G	A	A	T	4	9
22	A/G	G/A	A	A	T/C	G	A	A	T	13	8
23	A/G	G/A	A	A	T	G	A	A/G	T	13	10
24	G	A	A	A	T/C	G	A	A	T	4	8
25	G	A	A	A	T/C	G	A	A	T	9	8
26	G/A	A/G	A	A	T	G	A	A	T	4	13
27	G	A/G	A	A	T	G	A	A	T	9	1

The haplotype pairs shown in Table 4 were estimated from the unphased genotypes using an extension of Clark's algorithm (Clark, A.G. (1990) *Mol Bio Evol* 7, 111-122), as described in U.S. Provisional Application Serial No. 60/198,340 entitled "A Method and System for Determining Haplotypes from a Collection of Polymorphisms". In this method, haplotypes are assigned directly from individuals who are homozygous at all sites or heterozygous at no more than one of the variable sites. In our analysis, the list of haplotypes is augmented with haplotypes obtained from a three-generation Caucasian family and a two-generation African-American family. This list of haplotypes is then used to deconvolute the unphased genotypes in the remaining (multiply heterozygous) individuals.

By following this protocol, it was determined that the Index Repository examined herein and, by extension, the general population contains the 20 human NPY haplotypes shown in Table 5 below.

Table 5. Haplotypes Identified in the NPY Gene

Haplotype Number	PS 1	PS 2	PS 3	PS 4	PS 5	PS 6	PS 7	PS 8	PS 9	PS 10	PS 11	PS 12	PS 13	PS 14	PS 15	PS 16	PS 17	PS 18	PS 19
1	G	C	G	C	G	C	G	G	T	T	G	G	A	A	T	G	A	A	T
2	T	C	G	A	G	C	T	G	T	C	G	A	A	A	T	G	A	A	T
3	T	C	G	C	G	C	T	A	T	C	G	A	A	A	T	G	A	A	T
4	T	C	G	C	G	C	T	G	T	C	G	A	A	C	T	G	A	A	T
5	T	C	G	C	G	C	T	G	T	C	G	A	G	A	T	G	A	A	T
6	T	C	G	C	G	C	T	G	T	C	G	A	G	A	T	G	A	A	T
7	T	T	G	C	G	A	G	G	T	T	A	G	A	A	T	G	A	A	T
8	T	T	G	C	G	C	G	G	T	C	G	A	A	A	C	G	A	A	T
9	T	T	G	C	G	C	G	G	T	C	G	A	A	A	T	G	A	A	T
10	T	T	G	C	G	C	G	G	T	C	G	A	A	A	T	G	A	G	T
11	T	T	G	C	G	C	G	G	T	C	G	A	A	A	T	G	G	A	T
12	T	T	G	C	G	C	G	G	T	T	A	G	A	A	T	G	A	A	C
13	T	T	G	C	G	C	G	G	T	T	A	G	A	A	T	G	A	A	T
14	T	T	G	C	G	C	G	G	T	T	A	G	A	A	T	G	A	G	T
15	T	T	G	C	G	C	G	G	T	T	A	G	A	A	T	G	G	A	T
16	T	T	G	C	G	C	G	G	T	T	G	G	A	A	T	A	A	A	C
17	T	T	G	C	G	C	G	G	T	T	G	G	A	A	T	A	G	A	T
18	T	T	G	C	G	C	T	G	C	C	G	A	A	A	T	G	A	A	T
19	T	T	G	C	T	C	G	G	T	T	A	G	A	A	T	G	A	A	C
20	T	T	T	C	G	C	G	G	T	T	A	G	A	A	T	G	A	A	T

In view of the above, it will be seen that the several advantages of the invention are achieved and other advantageous results attained.

As various changes could be made in the above methods and compositions without departing from the scope of the invention, it is intended that all matter contained in the above description and shown in the accompanying drawings shall be interpreted as illustrative and not in a limiting sense.

All references cited in this specification, including patents and patent applications, are hereby incorporated in their entirety by reference. The discussion of references herein is intended merely to summarize the assertions made by their authors and no admission is made that any reference constitutes prior art. Applicants reserve the right to challenge the accuracy and pertinency of the cited references.

What is Claimed is:

1. A method for genotyping the neuropeptide Y (NPY) gene of an individual, comprising determining for the two copies of the NPY gene present in the individual the identity of the nucleotide pair at one or more polymorphic sites selected from the group consisting of PS1-PS5, PS9, PS13 and PS14.

2. The method of claim 1, wherein the determining step comprises:

   (a) isolating from the individual a nucleic acid mixture comprising both copies of the NPY gene, or a fragment thereof, that are present in the individual;

   (b) amplifying from the nucleic acid mixture a target region containing at least one of the polymorphic sites; .

   (c) hybridizing a primer extension oligonucleotide to one allele of the amplified target region;

   (d) performing a nucleic acid-template-dependent, primer extension reaction on the hybridized genotyping oligonucleotide in the presence of at least two different terminators of the reaction, wherein said terminators are complementary to the alternative nucleotides present at the polymorphic site; and

   (e) detecting the presence and identity of the terminator in the extended genotyping oligonucleotide.

3. A method for haplotyping the neuropeptide Y (NPY) gene of an individual which comprises determining, for one copy of the NPY gene present in the individual, the identity of the nucleotide at one or more polymorphic sites selected from the group consisting of PS1-PS5, PS9, PS13 and PS14.

4. The method of claim 3, wherein the determining step comprises

   (a) isolating from the individual a nucleic acid molecule containing only one of the two copies of the NPY gene, or a fragment thereof, that is present in the individual;

   (b) amplifying from the nucleic acid molecule a target region containing at least one of the polymorphic sites;

   (c) hybridizing a primer extension oligonucleotide to one allele of the amplified target region;

   (d) performing a nucleic acid template-dependent, primer extension reaction on the hybridized genotyping oligonucleotide in the presence of at least two different terminators of the reaction, wherein said terminators are complementary to the alternative nucleotides present at the polymorphic site; and

   (e) detecting the presence and identity of the terminator in the extended genotyping oligonucleotide.

5. A method for predicting a haplotype pair for the neuropeptide Y (NPY) gene of an individual comprising:

   (a) identifying an NPY genotype for the individual at two or more of the polymorphic sites PS1-PS5, PS9, PS13 and PS14;

    (b)   enumerating all possible haplotype pairs which are consistent with the genotype;

    (c)   accessing data containing the NPY haplotype pairs determined in a reference population; and

    (d)   assigning a haplotype pair to the individual that is consistent with the data.

6.   A method for identifying an association between a trait and at least one genotype or haplotype of the neuropeptide Y (NPY) gene which comprises comparing the frequency of the genotype or haplotype in a population exhibiting the trait with the frequency of the genotype or haplotype in a reference population, wherein the genotype or haplotype comprises a nucleotide pair or nucleotide located at one or more polymorphic sites selected from the group consisting of PS1-PS5, PS9, PS13 and PS14, wherein a higher frequency of the genotype or haplotype in the trait population than in the reference population indicates the trait is associated with the genotype or haplotype.

7   The method of claim 6, wherein the haplotype is selected from haplotype numbers 1-20 shown in Table 5.

8.   The method of claim 7, wherein the trait is a clinical response to a drug targeting NPY.

9.   A composition comprising at least one genotyping oligonucleotide for detecting a polymorphism in the neuropeptide Y (NPY) gene at a polymorphic site selected from PS1-PS5, PS9, PS13 and PS14.

10.   The composition of claim 9, wherein the genotyping oligonucleotide is an allele-specific oligonucleotide that specifically hybridizes to an allele of the NPY gene at a region containing the polymorphic site.

11.   The composition of claim 10, wherein the allele-specific oligonucleotide comprises a nucleotide sequence selected from the group consisting of of SEQ ID NOS:4-19, the complements of SEQ ID NOS: 4-19, and SEQ ID NOS:20-51.

12.   The composition of claim 9, wherein the genotyping oligonucleotide is a primer-extension oligonucleotide.

13.   An isolated polynucleotide comprising a nucleotide sequence selected from the group consisting of:

    (a)   a first nucleotide sequence which is a polymorphic variant of a reference sequence for neuropeptide Y (NPY) gene or a fragment thereof, wherein the reference sequence comprises SEQ ID NO:1 and the polymorphic variant comprises at least one polymorphism selected from the group consisting of guanine at PS1, cytosine at PS2, thymine at PS3, adenine at PS4, thymine at PS5, cytosine at PS9, guanine at PS13 and cytosine at PS14; and

    (b)   a second nucleotide sequence which is complementary to the first nucleotide sequence.

14.   The isolated polynucleotide of claim 13 which comprises a NPY isogene.

15. The isolated polynucleotide of claim 13 which is a DNA molecule and comprises both the first and second nucleotide sequences and further comprises expression regulatory elements operably linked to the first nucleotide sequence.

16. A recombinant nonhuman organism transformed or transfected with the isolated polynucleotide of claim 13, wherein the organism expresses a NPY protein encoded by the first nucleotide · sequence.

17. The recombinant organism of claim 16 which is a nonhuman transgenic animal.

18. The isolated polynucleotide of claim 13, wherein the first nucleotide sequence is a polymorphic variant of a fragment of the NPY gene, the fragment comprising one or more polymorphisms selected from the group consisting of guanine at PS1, cytosine at PS2, thymine at PS3, adenine at PS4, thymine at PS5, cytosine at PS9, guanine at PS13 and cytosine at PS14.

19. A computer system for storing and analyzing polymorphism data for the neuropeptide Y gene, comprising:

    (a) a central processing unit (CPU);

    (b) a communication interface;

    (c) a display device;

    (d) an input device; and

    (e) a database containing the polymorphism data;

wherein the polymorphism data comprises the genotypes and haplotype pairs shown in Table 4 and the haplotypes shown in Table 5.

20. A genome anthology for the neuropeptide Y (NPY) gene which comprises NPY isogenes defined by any one of haplotypes 1-20 shown in Table 5.

5

POLYMORPHISMS IN THE NPY GENE

```
GAGAGAAGAG TGGACACGTT TAGATACGGA AGATATAAAA AAGATTCATT
GCTTTATATT GTCCTGTAAC TTAAAAAAAA GATTCAAGTT GAAACTCTAG    152300
AGATGAAAGC TACCAGGTAT AAGGTGCAAA ATACACTGGA TAGGATTGGC
AGTAATCAGA TATTACAGAA GAAAACATTA GTGAACTCAC AGACATAGCA    152400
ATAGAGACTA TCCAAAATAA AATATGAAGG AAAAAATGAG CAAAACAGAA
GCAAAAACAG ATCACCAACA AACTGTGGGC CAATATACAC TCACCGTCTG    152500
ACTCAGCCAT TCTGGTTTTA AGTATTTTCC AAGAGAATTA AAGGCTTTAT
GTCCATACAA AGACTTGCAT ATGAATGTCA AGGCAGCTTT ATTCGTAAAA    152600
GTTTCAAGTG AAAATAACCA AAAATTCATG AGCAGTAGAA TTGATTAAAA
ATTTTGAAGT ATATCATATA ATGAAATACT GCCCAGCAAT AAAAAAGAAT    152700
AAACTATTGC TACATTTAAC AATGTGAGTG GATTCAAAAA TAATGATGCT
AAGTGAAAGA AGTCAGACAA AGAGTACCTA CCGTGGAACT ATATTTATGT    152800
GCAATTCTAG AAAACGCAAA CAAATCTCTA GTGATAGCAG ATCAATGGAG
GAGGGGGTGA GGAAGAGGCA AAAGGAAGGA ATTACAAAGG TACATGTGGA    152900
AATTTGGGGA AGTGATGTAT AATCTTCATT TTCTTGGTTT CAGTGATGGC
TTCCTGAGCA TATAACACAC ATGCATATGT GTAAGAATAT ATCCAATTGA    153000
ATATTTTACA CATTTGTAGC TTTGTATATA TTCAGCAGTA TCTATTCATT
CCTTCTGCTT CTTTACCTTA GGTTTAGTCA TGAAACCACT GGGTTTCATG    153100
TGGTCCAGGT TCTATCTGAA TCAAGCTATG AACCTGCTAT CAGTAAGACC
TCCTAATTGC AAACCACATG GACCTTTACC GGTTATTTGA CTTCTCTCGG    153200
TCATTACCTT CTCTAGTCTC CACTGATTCA CCTCCTTTTG AATCTCAACA
AGCTCAACAG TATCTCAATG TTATCTTAAC AAAGCTGTTA GAAATAAACA    153300
ACTTGACTTT GAAACAAAGG AGAGAAATGG CTCAGCAGCT AGGAGAAGAT
GCAGGGTTGA GTTGCCAACA GTTGGTTTTG TTTGTATTTT GGCCAGGGGA    153400
TGTGGCTTGG ACTGGAGAGA AAGGAGATAA GGATGTAAGC ACATGTAGGG
CATATCACCC CCTATTTTTT ATTCTCTGAA TCCTTAACCC TCAGAATAAG    153500
TTCTTATTCT TGAGAATCAA TGACATTATC TTAAGCTAAA TTAATCAAGC
CTCCACAGTG TTCTTCTCTC AATAGTGGTG TGGGCCTTCC TAGAAGTAAT    153600
TTTTCCCAAA TTCAGTGATA CATTTTAAGT TCAGATTTTA ATTGATATGA
ATCTGTGATA CACTCTAAAA TAAGATTATT TTATTGAAAA GTGGACTGTA    153700
ACTTTCCCTT TATCTAGGAA GAGCTCTAAG TTAGAAGATG TTTTGCACTT
TTACCGAAGG CTGTGTCTTG TAAGCACCCC CGAGCAACTC TGAGAGCCTT    153800
GATTTTTGTG TCCTCAGCAT ATGTTTGTGT AATACAGAAA GAGAAGCAGT
TGCCAAGTGA AAGGGATGTT GGTCTCCAAA ATTATAGTTT GATCCCACAA    153900
ACACACAAAC ACATACATGC AAAGGATTGT TTGCTTCACG GTTTTTGATA
TTTAATTCAA TGCTGTTGGA ACAGCACAAA AACTAAGTGT CAGTTTAACA    154000
GAATCACTTG TCCTTTTAGC ATTAAAATAA CATGGAACTT AATGCTTTAA
TTTCCCAACA TGCCTTTTTA TTTAGAAAGA TTCAGACTTA TATTTCATTT    154100
AGAAATAAAA TGCCATTTTA TTTAGAAAGA TACAGGAGCA TTCATTCACG
GAACTTTCAG ATCTCAGTCC ACTGCATAAA ATCTTGATCC TGTAATAATA    154200
GTTTCTGTAT CTTGCATATT CATTCAACAG GTTTAACGCG ATGAGCAAAT
                              G
TAATGTTCAT CGTTTTTAAC ATGTTTCATC TTAATCAGAA CCCACATTCT    154300
CAACGTTAAT TGAACGTACA TAGGACTATA CAAGGGTTAG TAAATAAGAC
AGAAACTGTT GTTCATTTAA CCACCGTCAC TTTGGACCAA AAAAGAAAAA    154400
ATATATATTT TTAAAATTGA GCTTAAAAGA GTCTCTAGAA GCTGGAAGCG
TGGCTCTTTT TCAGCAAACT GGGGGAATAG GTTTACCGTG TTCCCCCTCT    154500
GGGGAATTTT GAGTCGCCAC ACTCATGTCT CGACCGAGCC TGGCTCGCTG
CGTCTGAGCG AGTACTTGAG GAAGGCTGAT CTAGAAAAAC CAGCTGAGAG    154600
AAGGGGCAGA AGCCCCTGAA ACCACGGGCG GGGGTGGGGT GGGGAGCGCA
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FIGURE 1A

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GCTTTGGGAC CCTCTAGCCG GAGACTTCCG GCAGCTGCCT CCGACTTGTT        154700
CTAAGTACAG GAAAAATCTG TGCGCCCAGT TGCCTCACTC CAACAGCGCG
CAGTTGTGCC CGGCGAGGAT GCCGCGCTAG TCGTGGAGAT GCCCCACCAC        154800


AAAGAGGATT CAGGTGCTTC CTACTCCGGC ACCCAGTGGG TTGGTAGTCC
                                                 C
TGTTGGCAGG AGACAAGAAT CGTCTGGGCT GCTCCTATCT CTGGCAGGAC        154900
TAGACGGGGC GTGAAGGAAA GAAGGAAAGA AGGAAAGCAG GGATCGGGCA
CTGCCCGAGG GCAGATACTT GGGCTTTGGT GTTGTCCAGC GCGCTCGGAG        155000
TGCGCTGCCT CGCTCACGCG GTCCCAGGCC CCGCTTCTTC AGGCAGTGCC
TGGGGCGGGA GGGTTGGGGT GTGGGTGGCT CCCTAAGTCG ACACTCGTGC        155100
           T
GGCTGCGGTT CCAGCCCCCT CCCCCCGCCA CTCAGGGGCG GGAAGTGGCG
              A
GGTGGGAGTC ACCCAAGCGT GACTGCCCGA GGCCCCTCCT GCCGCGGCGA        155200
GGAAGCTCCA TAAAAGCCCT GTCGCGACCC GCTCTCTGCA CCCCATCCGC
TGGCTCTCAC CCCTCGGAGA CGCTCGCCCG ACAGCATAGT ACTTGCCGCC        155300
CAGCCACGCC CGCGCGCCAG CCACCGTGAG TGCTACGACC CGTCTGTCTA
GGGGTGGGAG CGAACGGGGC GCCCGCGAAC TTGCTAGAGA CGCAGCCTCC        155400
CGCTCTGTGG AGCCCTGGGG CCCTGGGATG ATCGCGCTCC ACTCCCCAGC
           TA.       T
GGACTATGCC GGCTCCGCGC CCCGACGCGG ACCAGCCCTC TTGGCGGCTA        155500
AATTCCACTT GTTCCTCTGC TCCCCTCTGA TTGTCCACGG CCCTTCTCCC
GGGCCCTTCC CGCTGGGCGG TTCTTCTGAG TTACCTTTTA GCAGATATGG        155600
           A
AGGGAGAACC CGGGACCGCT ATCCCAAGGC AGCTGGCGGT CTCCCTGCGG
GTCGCCGCCT TGAGGCCCAG GAAGCGGTGC GCGGTAGGAA GGTTTCCCCG        155700
GCAGCGCCAT CGAGTGAGGA ATCCCTGGAG CTCTAGAGCC CCGCGCCCTG
CCACCTCCCT GGATTCTTGG GCTCCAAATC TCTTTGGAGC AATTCTGGCC        155800
CAGGGAGCAA TTCTCTTTCC CCTTCCCCAC CGCAGTCGTC ACCCCGAGGT
GATCTCTGCT GTCAGCGTTG ATCCCCTGAA GCTAGGCAGA CCAGAAGTAA        155900
CAGAGAAGAA ACTTTTCTTC CCAGACAAGA GTTTGGGCAA GAAGGGAGAA
AAGTGACCCA GCAGGAAGAA CTTCCAATTC GGTTTTGAAT GCTAAACTGG        156000
   C
CGGGGCCCCC ACCTTGCACT CTCGCCGCGC GCTTCTTGGT CCCTGAGACT
           T          .              A
TCGAACGAAG TTGCGCGAAG TTTTCAGGTG GAGCAGAGGG GCAGGTCCCG        156100
ACCGGACGGC GCCCGGAGCC CGCAAGGTGG TGCTAGCCAG TCCTGGGTTC
G                    G                     .
TCTCTGCGGG ACTGGGACGA GAGCGGATTG GGGGTCGCGT GTGGTAGCAG        156200
GAGGAGGAGC GCGGGGGGCA GAGGAGGGAG GTGCTGCGCG TGGGTGCTCT
GAATCCCCAA GCCCGTCCGT TGAGCCTTCT GTGCCTGCAG ATGCTAGGTA        156300
   C
  [EXON 1: 156291
ACAAGCGACT GGGGCTGTCC GGACTGACCC TCGCCCTGTC CCTGCTCGTG
  C
TGCCTGGGTG CGCTGGCCGA GGCGTACCCC TCCAAGCCGG ACAACCCGGG        156400
              A
CGAGGACGCA CCAGCGGAGG ACATGGCCAG ATACTACTCG GCGCTGCGAC
ACTACATCAA CCTCATCACC AGGCAGAGGT GGGTGGGACC GCGGGACCGA        156500
             156478]
TTCCGGGAGC GCCAGTGCCT GCACACCAGG AGATCCTGGG GATGTTAGGG
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FIGURE 1B

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AAAGGGATTG TTTCTTTTCC TTCGCTCTAT CCCAGGGCAG GACAGTATCA        156600
GGCACTTAGT CAGCTCTAGG TAAATGTTTG TACAGGGCAC ACTCTACACA
AAATGGGTAC CTTCCATTTT GTGCAACTAC AGTCACAGAG TCGTGATCCC        156700
CAGATTCAGG TTCCCCAGGC TGGTAGGCTG GCAATCTCCT CTCACTCACC
TCTTATGGTT TGTTGTGGTT CTTACGGCAG TGGGGCCCGG TCCAGAAATC        156800
TCGAAAGTAC CCAGTGAAAG GGGCAAGAAT GCGCCAGAGA AATGCTGTAG
GGGGAAACGC TAGCAAGGTG TCTAGGAGAA ACAGAACGAC CACCAAAGAA        156900
AACCAAACCA AGGAGTAAAC TGCAGGGTTG CCACAGACAT TGTCAGACTT
TCCGGCCTGC CCAGGGCTAA TTGAATGACA GTATTTAGAA AAAGACAAAT        157000
AGAGCTATTT TCTTTTCTTC TTCACTGTGT TTTTTTTGCA GTCTCACTCC
CATATTTGAA AATTCTAGTC CCAATAATTA GAGATTTAGC CTGAAAACAG        157100
TTTATCTAAA ATGCAGAGGT CTCCCTATTT TATAGCTGGT GAATTCTGAA
AAGAAAGTGC CCACTTGGTA TTTCATTAAG AAGAAACTTA CGTTGGTCCA        157200
GTAACTTTAG GATGAATTCA GGGATTACAT GGAATCTGAG AAATTGTAAA        
CTTTATTCTG AACATTCCGA AAGTCAAAAT GAAATGCCAT ATTTTCACTC        157300
TCAACATTGA CAATTAAGGG AAAAATCAAA TCCAAACTAT TTTACAAATT
ACATAGGATT TAAATTTTGT TTTCAAAATT GGTTGATCTT AGTTCTAAAT        157400
CATTTAATAT CTTATTTTAA TTACTGAGAT TAAATGATTT TAGTCATTTG
ACCGTATCTA AAGGATAAGC CCTTTAGAAG CATTAATGTT AGTTAACATG        157500
TTTAAATTAT ATGTTTAAAG TATTGTGGGG TTTATATATT TATTTATTTA
GTCCATATTT TTACTGTTTA CAAGTCAGGG AGTAGTTGGA AAGGATCCAA        157600
AAGGGGGGAA GATTTATACT TTTAAAATAA AGTTTGCTTT TTCCTGTTTC
ATCAAAAAGG TATTTAACAT CCTAAAACAC TTTTGAAAAA GCCCCAATTT        157700
ATATAGAATC TGAAAAACAT GAAAATGCCA GAATTAAATA GTAGTTGTTT
GTGATCCCTT GAAATCTGAT CATTTTAATT AGCAAAAAGC AATAATATTT        157800
TATATTCCAT ACTTCCCAGA AAGTTAATTG ATTCAAATTC TTGCTATGTT
ATGCCGTCAT GCAGATCAAA GAGAACTTGT ACAAACAAAA AGATATTAAC        157900
TGTTTTTTCT AGAAAAACCT TTGTTAGTTA TTTAGCTGTA GCATAATACA
CAGTAGAACT CATTTTTCCG AAGATGATTG CCATAAAGCT TTACCATTTT        158000
ACAAAGCTCT AGCATTTTAA AATGTTTTCC TAGGCATTCT CTCATTCAGT
CGTTACCTCA AAGCTTTGAA GGGAGTATAA TTATCTTCAT TTCACAAATC        158100
ATGAAGCTAA TGACATTTCA TGGAAGCTAT TTTTATAGGA ATTATGGAAT
TGGAGCCCTC CTTTGAGTGT CTTTGGATGT TGTCACCACC ATCAACAGGC        158200
AGTGTTATCT CCAACATGTT TGATGAGAAA AAAACATAAA GGAAAGGGAC
TTGGGTTTCA TGGCAAGAGA TTCAGCTTAG ATGGGAATGA GCTGGTCTCT        158300
GAGGCAGAGT TTCCTGTCTC ATCTTTGTAT AAACTCCTTC TTAGAAACCA
GCAGAGGCTC TTAATGTACG TCCATCTTCT CCTTCCCACC ACAGACACCC        158400
ACTCCTATGT GCGTTTCTGA AAATTACAGG GTCATTGTG GATGTGCTTT
AATTTAGTAT AGATATAATC TGATATCATT ATCTTTCTTT CCCCCCGCCC        158500
CCCGCCACCA GCATTCCCCC TGCACTCAAT TTGTGCTTAT ATTCCTTTTG
CTGTATATAT ACCCCTGGCT CATTGCTTCT GGAAGCTGCA TTACTTATTC        158600
TAGAGTGTGG GTCTCCTACA ATTTATTTAT ATAGCCCCCT AAATCAGTGG
TTCTCAACTG GGGGTGATTT TATGCTCCCC AGGTGTGATT TGGCAGTGTC        158700
TGGAGACATT TTTGGTTTTC ACAACTGGAG GTGGGCACTG CTGGTACCTA
GGTGTATAGA TGCCAAGGAG GCTGCTAAAT ATCCTACCAT GCCCAGGACA        158800
GCCCCCATTA CACAGAATAG CCAGTCCCAG AGAAACTCAG CCTTAGATTA
GTGTTTCTCC TTCCTGGCTG CACATTAAAA TTATCAGGGG AGCTTTTAAA        158900
ATACACTGAT GCTGGGATCC ACCCCTGACC AATTAAATCA GACCCTCTGG
GATGGGGCCT GGGCTTTAGA GTGTGTTTTT AACACTCCCC AGGTGATTCT        159000
AATGTGCAGC CAGGTTTGAG AATCTTTACT CTAAGGACTC AATCTTTGAA
AGTTACAGCA TTGTAGAGGG GAAGGATGAT CCAGAATTAA TTAAAGAAGG        159100
AAGAAGAGTG AACAGTACTG AACTGGATGC AAGATTATGA CTGTGCAAAT
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FIGURE 1C

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CCTCCTATTT CAGATTTAGG TTCTTTACTG GTAAATTGGA TAAAAAGTTG    159200
GCAGTTCTTT TACTCACTGG AGCATGATGA AGATAAATGA GTTACAGTTG
CCCTAGACTT TTACGATTAG TAAAACCAAT GGATGGACAG AATAAATAAG    159300
TATAAGCAGA CACCTATAGT CTAATATCGC AACTCCAGAA AATCTTTGAG
AAAATCTCAC ACATGATGGG AGTGGGACAG GGAGTTGTTT TGTTTTTGCT    159400
TCCCTTGAGT CTCAGCAAAA GAACAGTCAT AGAAAATATT TCCAAAAAAT
TTAGATATAA TTTTGCTTGG CAATTAGAAT ATACCCTCTG TCAGCTTTGT    159500
TCAGTTGTTG GACCTATTAA CCAATCCATA CCAGACTAAT TTATTTCTAA
ACAAAGTGGC GAATGGTGTC CAATGCCTTT TCTGAGAAAG AGAAAAATGG    159600
CCTCTTCAAA AGACGCAATT TGGTATAGAC TTGTCTGGCC GTTTTGTAGT
GTTCTGGAGT GGGGAGCATA TACTACCCTC ACAGGTAAAC GAAACCTTAC    159700
CTGTATGTAT TGTCTCACCC ATATCCTGCT ACAGGAGAAA AAAAAAAAAA
AAAAAAAAG AACATTTCCA TGGATCAGTT AGATTCCTTT AGACTTACAG     159800
GGAGGCTAAA ATCACAGCAT TAGGAAAGAC TTTAGCATTT GGCACCTTGA
ATGAATATGG AAAATATAAG ACCCAGGGTC TGTTGGTAGC CAGAAGAATG    159900
GATGCATTTC AACAGCAGTA ATTTGCATGC GTGTATCACT TTGTCACTTT
GAAAGCATTG TGACTCACAT TTCACACTGG ATGTTCACAA AGACTCTGAT    160000
TTAAGCAGGG GTTGTTTTAC AAACGAGAAA ATTGGGATTC AAAGACACAG
GTGGGTCTTG GACGGCAATC TGCTGATTTT TTTTTTCTAG GACTGTCTTT    160100
GAAATATCAT CTGGTCTCCT TGTTAAGATT TAAACTGCTG TGTATCATCA
GAACAATAGC CAATTAACTA ATTGTGGAAA CATTTACAAT CATGCTTCCA    160200
TGTTTTCAAA TTTCCTGTCA CCTAGTTATA AAAGTATGTA TCTCTATACA
TATTATTTGT GGTTGTTTAA GAAGGCAGGA ATTTGACTAG GAATTTTGGT    160300
TCGGTTTAAA TAACCTCCTA GATTATTTAT TATTTGAGCA AAGTTTAAAT
ATATATACAA ATAATTTGAA ATAATATTTA TGATAATGTT TCGATAATAT    160400
ATGCTTCATA CACCTAGCTT GCTTGTTACA GATGAACACC TGACAATAAT
GTTTAGTTTT CATATCCCAA ATAGGAGACT ATCTTCCTTT TTCCTAAAGG    160500
TTTTTATGCC TATTCCAAAC TTGCTTTAAA AGACTTTTT TTTTCCAGAT
    [EXON 2: 160549.
ATGGAAAACG ATCCAGCCCA GAGACACTGA TTTCAGACCT CTTGATGAGA    160600
GAAAGCACAG AAAATGTTCC CAGAACTCGG TATGACAAGG CTTGTGATGG
        160629]
GGACATTGTT GCAGAGCTCA AGGTGCCCAG GGGAGGGAAG TCAGAGACAG    160700
GTGCCTGGTG GGAGGCACCA CCAGGCTTCT AGACTAGGGG AGATTTCGGC
AGAAATGAGG ATGAAGAAGC AGGCAGAGGA GACCTCTCTT TTCATGTACT    160800
CATTGTCAAA TCTCTAAACT TCTGTGGAAT AACATTTTTT TAGCTCAGGT
TATTTGGTGC TCCTTTGACG ATTTTAAAAA TAATTGACTT TTAACTTAGA    160900
AATTATATAG AATTTCTAAC ATTGTTGGAG TCAGGGGAGC ATAGAGTGAG
TCCCTGACTC TAAATTGAAA TGACCAACTG TCCATCTCTG GCTGGGAAGA    161000
TCTGTAGATT TTATATATGT GGAGTGTCCA GATGTCCTAG GAAATGTCCC
ATGTGGTTTT TACTCCAACT TTCCCAGTTT TACATGAAGC TGTAGGTAAA    161100
TTAGGTGCTC TGGATTACTT TTTGACATAT TTTAATTCAA CTTGTGTCAT
ATAGTGATTT ATTACATTCA ACGAATGCAC GTTGAATGAC AGTATTCTTA    161200
GGAGTTTATC TGTAGTCTCA GGACTTTCTT GCAGTTTGGG TCTGTCTTGG
AAAGAGGTGC CCACCAATGT CTAAGTGTTA AACATTGAGC TAAGGTTTTC    161300
TGGTGCCTCC TGGACAAGGA GTGCTGCTGA AGAGGAGCCT GGTCCTCTTC
CATCTTCATA ACCTCCTGAC TCTTCAGTCC TTGCTTCCTG TCTCTAACCA    161400
CAGTAAGATC ATAGTTCACA GGGTCCACCC ACGTCTGTGG CAGCTTGGCT
GCCTTAAACC AAGTTCTACA ATGTCAAGAA CCAAGTTCCC CAGAAGAGGT    161500
CACTACAACA CTTGCTCAGT GAGTGTGACT GACGTGGCTC TCCATCAAGT
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FIGURE 1D

```
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GTCTCCTTCT GTCACTCAGG CCAGAGCACA GTGGTGCAAT CAAAGATCAC        161900
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GTTTGCCACA GTAAAAAGTT TAGGGTAGAC TATATCGACT TACTGTTAGA        162100
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FIGURE 1E

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FIGURE 1F

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CCCAGGTACT TGGGAGGCTG AGACAGGAGA ATCACTT                  166737
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FIGURE 1G

POLYMORPHISMS IN THE CODING SEQUENCE OF NPY

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GCGCTGCGAC ACTACATCAA CCTCATCACC AGGCAGAGAT ATGGAAAACG        200
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AAAATGTTCC CAGAACTCGG CTTGAAGACC CTGCAATGTG GTGA              294
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FIGURE 2

ISOFORMS OF THE NPY PROTEIN

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FIGURE 3

SEQUENCE LISTING

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    Nandabalan, Krishnan
    Stephens, J. Claiborne
    Chew, Anne
    Denton, R. Rex
    Lanz, Elizabeth M.

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A.	**CLASSIFICATION OF SUBJECT MATTER**
IPC(7)	:C07H 21/02; C12Q 1/68; C12P 21/00, 19/34; A01N 43/04; G06F 17/00
US CL	:Please See Extra Sheet.

According to International Patent Classification (IPC) or to both national classification and IPC

B.	**FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 536/23.1, 24.3, 24.32, 24.33; 436/6, 91.1, 91.2;800/4; 514/44; 345/418

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

Medline, West 2.0, Embase, Scisearch, Biotechds, Caplus, Canadian Patents

C.	**DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,602,024 A (GERALD et al) 11 February 1997. See entire patent, especially figure 5.	13-15
X	US 5,985,616 A (PARKER et al.) 16 November 1999. See entire patent, especially SEQ ID NO: 2.	13-15
A, P	WO 00/18960 A2 (MASSACUSETTS INSTITUTE OF TECHNOLOGY) 06 April 2000. See entire reference.	4

☐ Further documents are listed in the continuation of Box C.   ☐ See patent family annex.

*	Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance		
"E"	earlier document published on or after the international filing date	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O"	document referring to an oral disclosure, use, exhibition or other means		
"P"	document published prior to the international filing date but later than the priority date claimed	"&"	document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
29 MARCH 2001	**24 APR 2001**
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231· Facsimile No. (703) 305-3230	Authorized officer CYNTHIA B. WILDER Telephone No. (703) 308-0196

Form PCT/ISA/210 (second sheet) (July 1998)★

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US00/34758

Box I   Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
   because they relate to subject·matter not required to be searched by this Authority, namely:

2. ☒ Claims Nos.: 1-4, 9-12, 18 and PS1
   because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

   Please See Extra Sheet.

3. ☐ Claims Nos.:
   because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II   Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

   Please See Extra Sheet.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
   1-4, 9-15, 18 and PS1

Remark on Protest   ☐ The additional search fees were accompanied by the applicant's protest.
                        ☐ No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (continuation of first sheet(1)) (July 1998)★

A. CLASSIFICATION OF SUBJECT MATTER:
US CL :

536/23.1, 24.3, 24.32, 24.33; 436/6, 91.1, 91.2;800/4; 514/44; 345/418

BOX I. OBSERVATIONS WHERE CLAIMS WERE FOUND UNSEARCHABLE
2. Where no meaningful search could be carried out, specifically:

Claims 1-4, 9-12, 18 and PS1 could not be searched because the claims are drawn to a composition and method of genotyping comprising the polymorphic site 1 wherein a G is substituted for a T at position 154224 of the NPY gene in figure 1 or Genbank nucleotide position 31293 of the NPY gene. The NPY gene reference sequence as depicted as SEQ ID NO: 1 only has 14537 nucleotide bases and there are no positions corresponding to 154224 or 31293 as recited in Table 3 of the description (page 30). Additionally figure 1 is not represented by a sequence identifier (SEQ ID NO:). Therefore a meaningful search of the positions as claimed in claims 1-4, 9-12, 18 and PS1 cannot be obtained. Because claims 13-15 read on a fragment of the reference NPY gene sequence, a search was performed on those claims.

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING
This ISA found multiple inventions as follows:

This application contains the following inventions or groups of inventions which are not so linked as to form a single inventive concept under PCT Rule 13.1. In order for all inventions to be searched, the appropriate additional search fees must be paid.

Groups 1-8, claim(s)1-4, 9-15 and 18, in part, drawn to method of genotyping the neuropeptide Y gene of an individual with one or more polymorphic sites (PS1-PS5, PS9, PS13, PS14), an isolated polynucleotide comprising a nucleotide sequence with a polymorphic variant for the NPY gene or a fragment thereof, and a composition comprising using one of the 8 polymorphic sites, respectively. For example, if group 1 is elected, the claims 1-4, 9-15 and 18 will be examined to the extend that they apply to PS1. If group 8 is elected, the claims 1-4, 9-15 and 18 will be examined to the extend that they apply to PS14.

Groups 9-40, claim(s) 5, in part, drawn a method for predicting the haplotype pair for the NPY gene by identifying an NPY genotype for the individual at two or more polymorphic sites PS1-PS5, PS9, PS13, PS14. For example if group I is elected , the claim 5 will be examined to the extend that it applies to PS1 and PS2. If the group 40 is elected, the claim 5 will be examined to the extend that it applies to PS13 and PS14.

Groups 41-48, claim(s) 6-8, in part, drawn to a method of identifying an association between a trait and a genotype or haplotype between one of the 8 PS sites of the NPY gene. For example, if group 41 is elected, the claims 6-8 will be examined to the extend that they apply to PS1.

Groups 49-56, claim(s)16-17, in part, drawn to a nonhuman recombinant organism comprising one of the 8 PS sites, respectively. If group 49 is elected, the claims 16-17 will be examined to the extend that they apply to PS1.

Groups 57-76, claim(s) 19-20, in part, drawn to a computer system and genome anthology (database) comprising the polymorphism data of one of the 20 haplotypes in Table 5, respectively. If group 57 is elected, the claims 19-20 will be examined to the extend that they apply to the first of the 20 haplotypes.

The inventions listed as Groups 1-76 do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The broadest recitation of the first claim product, namely a fragment of the nucleic acid of the neuropeptide Y gene (NPY)of claim 15, is known and is well characterized in the prior art. Thus the "special technical feature" is not special and is not a contribution over the prior art. Furthermore, Gerald et al. (5,602,024, 11 Feb 1997) teach fragments of the NPY gene. Parker et al. (5,985,616, 16 NOV 1999) teach fragments of the NPY gene. These fragments of the NPY gene thus meets the requirements of Groups 1-8. Hence there is not a special technical feature linking the groups.

The special technical feature of Groups 1, PS1 is lacking in groups 2-8, and 37-76. The nucleic acid of Groups 1 is not required in Group 2 such that the PS site is not required for the examination of PS 1 or visa versa. Additionally, the

different PS sites are structurally and functionally distinct as shown in Table 3 and requires different searches. Similar reasons can be set forth for the unique PS1 sites and unique pairs of PS1 sites. The non-human organisms (transgenic animals) are distinct products from the nucleic acid. Finally the computer system which requires data, does not rely upon the chemical structure of the nucleic acid or the non-human organism.

The groups (1-8), (9-40) and (41-48) are not so linked as a single inventive entity. The different groups are drawn to different methods requiring different starting reagents, different method steps and different objectives. Additionally, the method of genotying the NPY gene is not required or is necessary for predicting a haplotype pair or determining traits and visa versa.

Form PCT/ISA/210 (extra sheet) (July 1998)★